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OM protein - protein search, using sw model

Run on: March 20, 2000, 05:31:20 ; Search time 35.25 Seconds
(without alignments)
249.964 Million cell updates/sec

Title: US-08-509-359B-138
Perfect score: 1923
Sequence: 1 EELTKYGAHVIMLFVPVT.....STDNLVRPFMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 188963 seqs, 23686106 residues

Database : A_Geneseq_36.*

Word size : 0

Number of hits that pass the threshold : 188963

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1923	100.0	448	1 W05762	Human presenilin-2
2	1923	100.0	448	1 W11321	Human AD4 protein.
3	1923	100.0	448	1 W23967	Human presenilin-2
4	1919	99.8	448	1 W05763	Presenilin-2 M239V
5	1918	99.7	448	1 W05765	Presenilin-2 I420T
6	1914	99.5	448	1 W05764	Presenilin-2 N141I
7	1907.5	99.2	447	1 W28508	Full AD4/AD3LP seq
8	1723	89.6	414	1 W05766	Presenilin-2 delta
9	1591.5	82.8	376	1 W28506	AD4/AD3LP sequence
10	1431.5	74.4	467	1 W05735	Murine presenilin.
11	1431.5	74.4	467	1 W23966	Mouse presenilin-1
12	1429.5	74.3	467	1 W11839	Human early onset
13	1429.5	74.3	463	1 W11840	Early onset Alzhei
14	1429.5	74.3	467	1 W05733	Presenilin-1-1. Ne
15	1429.5	74.3	407	1 W28507	Partial AD3 sequen
16	1429.5	74.3	467	1 W11430	PS1/467 protein. D
17	1429.5	74.3	429	1 W41429	PS1/429 protein. D
18	1429.5	74.3	467	1 W23964	Human presenilin-1
19	1426.5	74.2	467	1 W05755	Presenilin-1-1 L28
20	1426.5	74.2	467	1 W05758	Presenilin-1-1 L39
21	1426.5	74.2	467	1 W05737	Presenilin-1-1 V82
22	1426.5	74.2	467	1 W05746	Presenilin-1-1 I21
23	1426.5	74.2	467	1 W41431	Mouse PS1/467 prot
24	1425.5	74.1	467	1 W05754	Presenilin-1-1 A28
25	1425.5	74.1	467	1 W05736	Presenilin-1-1 A79
26	1425.5	74.1	467	1 W05747	Presenilin-1-1 I23
27	1425.5	74.1	467	1 W05749	Presenilin-1-1 A26
28	1424.5	74.1	467	1 W12376	Human S182 gene pr
29	1424.5	74.1	467	1 W05738	Presenilin-1-1 V96
30	1424.5	74.1	467	1 W05739	Presenilin-1-1 Y11
31	1424.5	74.1	467	1 W05741	Presenilin-1-1 I14
32	1424.5	74.1	467	1 W05748	Presenilin-1-1 A24
33	1424.5	74.1	467	1 W27176	Human S182 gene, P
34	1423.5	74.0	467	1 W05753	Presenilin-1-1 E28
35	1423.5	74.0	467	1 W05757	Presenilin-1-1 G38
36	1423.5	74.0	463	1 W05734	Presenilin-1-2. Ne
37	1423.5	74.0	467	1 W05740	Presenilin-1-1 M13
38	1423.5	74.0	467	1 W05742	Presenilin-1-1 M14
39	1423.5	74.0	463	1 W23965	Human presenilin-1

40 1422.5 74.0 467 1 W05744 Presenilin-1-1 L17
41 1422.5 74.0 467 1 W56770 Homo sapiens PS-1.
42 1421.5 73.9 467 1 W05752 Presenilin-1-1 P26
43 1421.5 73.9 463 1 W42375 Human presenilin 1
44 1420.5 73.9 467 1 W05743 Presenilin-1-1 H16
45 1420.5 73.9 467 1 W05745 Presenilin-1-1 G20

ALIGNMENTS

RESULT 1

W05762

ID W05762 standard; Protein; 448 AA.

AC W05762;

DT 25-JUL-1997 (first entry)

DE Human presenilin-2.

KW Presenilin-2; human; hPS1-1; hPS1-2; PS-2; integral membrane protein; AD;

KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;

KW depression; antibody; gene expression modulator; therapy.

OS Homo sapiens.

PN W09634099-A2.

PD 31-OCT-1996.

PF 29-APR-1996; CA0263.

PR 28-APR-1995; US-431048.

PR 28-JUN-1995; US-496841.

PR 31-JUL-1995; US-509359.

PA (HSCR-) HSC RES & DEV LP.

PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.

PI Fraser PE, Rommens JM, St George-Hyslop PH;

DR WPI: 96-497631/49.

DR N-PSDB; T40031.

PT New presenilin genes - useful for diagnosis, therapy and drug

PT screening of familial Alzheimer's disease, cerebral disorders, etc.

PS Claim 4; Page 148-150; 178pp; English.

CC This sequence represents the wild type human presenilin-2.

CC W05733 and W05734 represent the two different forms of wild type human

CC presenilin-1 (PS-1). The form represented by W05734 results from

CC alternate splicing of the genomic DNA sequence. W05735 represents the

CC coding sequence for wild type mouse PS-1. The presenilins are a family of

CC highly conserved integral membrane proteins, with a common structural

CC motif, common alternate splicing patterns, and common mutational hot spot

CC regions. Mutations in PS genes are implicated in familial Alzheimer's

CC disease (AD) and possibly other diseases such as cerebral haemorrhage,

CC schizophrenia, depression etc., so detection of mutations in the DNA

CC encoding these sequences can be used for diagnosis of these diseases.

CC These proteins, or vectors that express them or containing antisense

CC sequences, antibodies selective for mutant forms of these proteins (such

CC as W05736) and modulators of PS gene expression are potentially useful

CC for treatment of AD etc. Transgenic animals are useful as models for drug

CC screening. The antibodies can also be used e.g. for affinity purification

CC and in immunoassays.

CC Sequence 448 AA;

SQ

Query Match 100.0%; Score 1923; DB 1; Length 448;

Best Local Similarity 100.0%; Pred. No. 1.6e-200;

Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 EELTKYGAHVIMLFVPVTLCHIVVVAIKSVRFYTERKNGQLIYPTFTDTPSGQRLL 60

|||||

DB 77 EELTKYGAHVIMLFVPVTLCHIVVVAIKSVRFYTERKNGQLIYPTFTDTPSGQRLL 136

|||||

QY 61 NSVLTLMISVTVVMTIFLWVLYKYRCYKFIHGWLIMSSLMFLFTYTYLGVVLKTYN 120

|||||

DB 137 NSVLTLMISVTVVMTIFLWVLYKYRCYKFIHGWLIMSSLMFLFTYTYLGVVLKTYN 196

|||||

QY 121 VAMDYPTLLITVYVNFAGVGVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAVIL 180

|||||

DB 197 VAMDYPTLLITVYVNFAGVGVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAVIL 256

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QY 181 GAISVYDVLVAVLCPKGPLRMLVETAQRNEPIFPALIYSSAMVYVTVGMALDPSQGAQLQ 240

|||||

Domain 225..244
/label= TM5
/note= "transmembrane domain 5"
245..249
/label= TM5-6
/note= "hydrophilic loop"
250..268
/label= TM6
/note= "transmembrane domain 6"
269..387
/label= TM6-7
/note= "hydrophilic loop"
388..409
/label= TM8
/note= "transmembrane domain 8"
Misc_difference 141
/note= "Asn141Ile mutation site (Claim 19)"
Misc_difference 239
/note= "Met239Val mutation site (Claim 19)"
Misc_difference 420
/note= "Ile420Thr mutation site"
WO9801549-A2.
PD 15-JAN-1998.
PF 04-JUL-1997; CA0475.
PR 02-JAN-1997; US-034590.
PR 05-JUL-1996; US-021673.
PR 12-JUL-1996; US-021700.
PR 08-NOV-1996; US-029895.
PA (HSCR-) HSC RES & DEV LP.
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
DR WPI: 98-286355/25.
DR N-PSDB: V04669.
PT New isolated mutant presenilin-1 genes - useful for developing
PT products for use in detection, diagnosis and therapy of Alzheimer's
PT disease and for drug screening
PS Claim 19: Page 203-204; 238pp; English.
CC This polypeptide comprises human presenilin-2 (hPS2). Its amino
CC acid sequence was deduced from an isolated cDNA clone (see V04669).
CC Human and murine presenilin-1 sequences are also provided (see
CC W23964-66). Mutations in the PS-1 and PS-2 genes are linked to
CC the development in humans of forms of familial Alzheimer's disease
CC (FAD) and may be causative of other disorders, e.g. cognitive,
CC intellectual, neurological or physiological disorders such as
CC cerebral haemorrhage, schizophrenia, depression, mental retardation
CC and epilepsy. Use of the nucleic acids and proteins comprising or
CC derived from the presenilins is made in screening and diagnosing
CC FAD, identifying and developing therapeutics for treatment of FAD,
CC and in producing cell lines and transgenic animals useful as models
CC of FAD. Methods for identifying substances that bind to, or
CC modulate the activity of a presenilin protein, and methods for
CC identifying substances that affect the interaction of a
CC presenilin-interacting protein with a presenilin protein are also
CC disclosed.
SQ Sequence 448 AA;

Query Match 100.0%; Score 1923; DB 1; Length 448;
Best Local Similarity 100.0%; Pred. No. 1.6e-200;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 EELTKYGAHVIMLFPVTLGMVAVATKSVRFYTERKNGQLIYPTFTDTPSVGQRL 60
DB 77 EELTKYGAHVIMLFPVTLGMVAVATKSVRFYTERKNGQLIYPTFTDTPSVGQRL 136
QY 61 NSVLNTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 120
DB 137 NSVLNTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 196
QY 121 VAMDYPTLLLTWNFGAVGVCVHKGKPLVLOQAYLIMISALMALVFYIKYLPWSAWVIL 180
DB 197 VAMDYPTLLLTWNFGAVGVCVHKGKPLVLOQAYLIMISALMALVFYIKYLPWSAWVIL 256

QY 181 GAISVYDLVAVLCPKGPLRMLVETAQERNEPIFPALYSSAMVWTVGMKLDPSOGALQ 240
DB 257 GAISVYDLVAVLCPKGPLRMLVETAQERNEPIFPALYSSAMVWTVGMKLDPSOGALQ 316
QY 241 LPYDPEMEEDSYDSFGPEPSYPEVPEPLTGYPGEELEEEERGKVLGLGDFIFYSVLVGK 300
DB 317 LPYDPEMEEDSYDSFGPEPSYPEVPEPLTGYPGEELEEEERGKVLGLGDFIFYSVLVGK 376
QY 301 AATGSGDWNNTLACFAVAILIGLCLTLLLLAVFKKALPALPISITFGLIYFSTDNLVRP 360
DB 377 AATGSGDWNNTLACFAVAILIGLCLTLLLLAVFKKALPALPISITFGLIYFSTDNLVRP 436
QY 361 FMDTLASHQLYI 372
DB 437 FMDTLASHQLYI 448

RESULT 4
W05763
ID W05763 standard; Protein; 448 AA.
AC W05763;
DT 25-JUL-1997 (first entry)
DE Presenilin-2 M239V mutation.
KW Presenilin-2; human; hPS1-1; hPS1-2; integral membrane protein; AD;
KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;
KW depression; antibody; gene expression modulator; therapy; muten.
OS Homo sapiens.
FH Key Location/Qualifiers
FT modified_site 239 /label= M239V
FT W09634099-A2.
PD 31-OCT-1996.
PF 29-APR-1996; CA0263.
PR 28-APR-1995; US-431048.
PR 28-JUN-1995; US-496841.
PR 31-JUL-1995; US-509359.
PA (HSCR-) HSC RES & DEV LP.
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
DR WPI: 96-497631/49.
PT New presenilin genes - useful for diagnosis, therapy and drug
PT screening of familial Alzheimer's disease, cerebral disorders, etc.
PS Claim 4: Page -; 178pp; English.
CC W05763-W05766 represent mutated versions of the human presenilin-2
CC protein (see W05762 for wild type sequence). The presenilins are a family
CC of highly conserved integral membrane proteins with a common structural
CC motif, common alternate splicing patterns, and common mutational hot spot
CC regions. Mutations in PS genes are implicated in familial Alzheimer's
CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
CC schizophrenia, depression etc., so detection of mutations in the DNA
CC encoding the wild type sequences can be used for diagnosis of these
CC diseases. The wild type proteins, or vectors that express them or
CC containing antisense sequences, antibodies selective for these mutant
CC forms of the proteins and modulators of PS gene expression are
CC potentially useful for treatment of AD etc. Transgenic animals are useful
CC as models for drug screening. The antibodies can also be used e.g. for
CC affinity purification and in immunoassays.
SQ Sequence 448 AA;

Query Match 99.8%; Score 1919; DB 1; Length 448;
Best Local Similarity 99.7%; Pred. No. 4.3e-200;
Matches 371; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 EELTKYGAHVIMLFPVTLGMVAVATKSVRFYTERKNGQLIYPTFTDTPSVGQRL 60
DB 77 EELTKYGAHVIMLFPVTLGMVAVATKSVRFYTERKNGQLIYPTFTDTPSVGQRL 136
QY 61 NSVLNTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 120
DB 137 NSVLNTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 196
QY 121 VAMDYPTLLLTWNFGAVGVCVHKGKPLVLOQAYLIMISALMALVFYIKYLPWSAWVIL 180


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QY 241 LPYDPEME-----EDSYDSFGSPSYPEVEFPPLTGYPG-----EEL----- 277
Db 309 VPKNPKYNTQRAERETQDSGNDGGFSEWEAQRDHLGPHRSTPESRAAVQELSGSI 368
QY 277 ---EBEERGKVLGDFIFYSVLVKGAAATGSGDNTTACFAVAILGICLTLLLAVF 333
Db 369 LTSEDPEERGKVLGDFIFYSVLVKGASATASGDWNTTIACFAVAILGICLTLLLAIF 428
QY 334 KKALPALPISITFGLIFFYSTDLNLRPSPMDTLASHQLYI 372
Db 429 KKALPALPISITFGLVFYFATDYLVOFPMDQLAFHFQYI 467

RESULT 11
W23966
ID W23966 standard; Protein; 467 AA.
AC W23966;
DE Mouse presenilin-1 (first entry)
KW Presenilin-1; Psi gene; mouse; familial Alzheimer's disease; FAD;
KW cerebral haemorrhage; schizophrenia; depression; epilepsy;
KW mental retardation; diagnosis; therapy; transgenic animal.
OS Mus musculus.
FH Key
FT Domain
FT 82..100
FT /label= TM1
FT /note= "transmembrane domain 1"
FT 101..132
FT /label= TM1-2
FT /note= "hydrophilic loop"
FT 133..154
FT /label= TM2
FT /note= "transmembrane domain 2"
FT 155..163
FT /label= TM2-3
FT /note= "hydrophilic loop"
FT 164..183
FT /label= TM3
FT /note= "transmembrane domain 3"
FT 184..194
FT /label= TM3-4
FT /note= "hydrophilic loop"
FT 195..212
FT /label= TM4
FT /note= "transmembrane domain 4"
FT 213..220
FT /label= TM4-5
FT /note= "hydrophilic loop"
FT 221..238
FT /label= TM5
FT /note= "transmembrane domain 5"
FT 239..243
FT /label= TM5-6
FT /note= "hydrophilic loop"
FT 244..262
FT /label= TM6
FT /note= "transmembrane domain 6"
FT 263..407
FT /label= TM6-7
FT /note= "hydrophilic loop"
FT 408..428
FT /label= TM8
FT /note= "transmembrane domain 8"
FT Misc_difference 177
FT /note= "Phel77Ser mutation site (Claim 1)"
FT Misc_difference 439
FT /note= "Ile439Val mutation site (Claim 1)"
FN W09801549-A2.
PD 15-JAN-1998.
PF 04-JUL-1997; CA0475.
PR 02-JAN-1997; US-034590.
PR 05-JUL-1996; US-021673.
PR 12-JUL-1996; US-021700.

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PR 08-NOV-1996; US-029895.
PA (HSCR-) HSC RES & DEV LP.
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
DR WPI; 98-286355/25.
DR N-PSDB; V04668.
PT New isolated mutant presenilin-1 genes - useful for developing
PT products for use in detection, diagnosis and therapy of Alzheimer's
PT disease and for drug screening
PT Disclosure; Page 199-200; 238pp; English.
CC This polypeptide comprises the murine presenilin-1 (PS1) homologue.
CC Its amino acid sequence was deduced from an isolated cDNA clone
CC (see V04668). Mutations in the human PS1 and PS2 genes (see
CC V04666-68) have been linked to the development in humans of forms
CC of familial Alzheimer's disease (FAD). All amino acids that are
CC mutated in analysed FAD pedigrees (see W23964) were conserved in
CC the murine homologue. Use of the nucleic acids and proteins
CC comprising or derived from presenilins can be made in screening and
CC diagnosing FAD, identifying and developing therapeutics for
CC treatment of FAD, and in producing cell lines and transgenic
CC animals useful as models of FAD. Methods for identifying
CC substances that bind to, or modulate the activity of a presenilin
CC protein, and methods for identifying substances that affect the
CC interaction of a presenilin-interacting protein with a presenilin
CC protein are also disclosed.
SQ Sequence 467 AA;

Query Match 74.4%; Score 1431.5; DB 1; Length 467;
Best Local Similarity 71.2%; Pred. No. 4e-147;
Matches 284; Conservative 36; Mismatches 50; Indels 29; Gaps 4;

QY 1 EELTLKYGAHVIMLPVPTLGMIVVVAIKSVRTEKNGQLIYPTFETDPSVGORLL 60
Db 71 EELTLKYGAHVIMLPVPTLGMIVVVAIKSVRTEKNGQLIYPTFETDPSVGORAL 130
QY 61 NSVLNTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSIMLLEFLTYIYLGVLKYN 120
Db 131 HSLNAAIMISIVIMTILLWLYKYRCYKVIHAWLIISLLLEFLTYIYLGVLKYN 190
QY 121 VAMDYPTLLTVMFGAVGVCVHMGVPLVQAYLIMISALMALVFIKYLPEWSAVIL 180
Db 191 VAVDYVTVALLIWNFGVGNIAIHMGVPLVQAYLIMISALMALVFIKYLPEWTAWLIL 250
QY 181 GAISVYDLVAVLCPKGPLRMLVETAQERNEPIFPALYISAMVYTMGMALDPSSOGALQ 240
Db 251 AVISVYDLVAVLCPKGPLRMLVETAQERNEPIFPALYISAMVYTMGMALDPSSOGALQ 308
QY 241 LPYDPEME-----EDSYDSFGSPSYPEVEFPPLTGYPG-----EEL----- 277
Db 309 VPKNPKYNTQRAERETQDSGNDGGFSEWEAQRDHLGPHRSTPESRAAVQELSGSI 368
QY 277 ---EBEERGKVLGDFIFYSVLVKGAAATGSGDNTTACFAVAILGICLTLLLAVF 333
Db 369 LTSEDPEERGKVLGDFIFYSVLVKGASATASGDWNTTIACFAVAILGICLTLLLAIF 428
QY 334 KKALPALPISITFGLIFFYSTDLNLRPSPMDTLASHQLYI 372
Db 429 KKALPALPISITFGLVFYFATDYLVOFPMDQLAFHFQYI 467

RESULT 12
W11839
ID W11839 standard; Protein; 467 AA.
AC W11839.
DT 07-MAY-1997 (first entry)
DE Human early onset Alzheimer's disease (EOAD) polypeptide.
KW Early onset Alzheimer's disease; EOAD; neurodegenerative disease;
KW diagnosis; therapy; inhibitor; antagonist; antibody.
OS Homo sapiens.
FH Key
FT Location/Qualifiers
FT misc_difference 26..29
FT /note= "unidentified amino acid residues"

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CC alternate splicing of the genomic DNA sequence. W05762 represents the
 CC coding sequence for wild type human PS-2. The presenilins are a family of
 CC highly conserved integral membrane proteins with a common structural
 CC motif, common alternate splicing patterns, and common mutational hot spot
 CC regions. Mutations in PS genes are implicated in familial Alzheimer's
 CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
 CC schizophrenia, depression etc., so detection of mutations in the DNA
 CC encoding these sequences can be used for diagnosis of these diseases.
 CC These proteins, or vectors that express them or containing antisense
 CC sequences, antibodies selective for mutant forms of these proteins (such
 CC as W05736) and modulators of PS gene expression are potentially useful
 CC for treatment of AD etc. Transgenic animals are useful as models for drug
 CC screening. The antibodies can also be used e.g. for affinity purification
 CC and in immunoassays.
 SQ Sequence 467 AA;

Query Match 74.3%; Score 1429.5; DB 1; Length 467;
 Best Local Similarity 72.3%; Pred. No. 6.6e-147;
 Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

Qy 1 EELTKYGAHVIMLFVPTLCMVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRLL 60
 Db 71 EELTKYGAHVIMLFVPTLCMVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRLL 130
 Qy 61 NSVLNTLMISVIIVMTIFLVLYKYRCYKFTGHWLIMSSLMLEFLTYIYLGEVLKTYN 120
 Db 131 HSLNAAIMISVIIVMTIFLVLYKYRCYKFTGHWLIMSSLMLEFLTYIYLGEVLKTYN 190
 Qy 121 VANDYITVALLIWNFGVGMISIHKGLPLRQQAAYLIMISALMALVFIKYLPEWTAWLIL 180
 Db 191 VANDYITVALLIWNFGVGMISIHKGLPLRQQAAYLIMISALMALVFIKYLPEWTAWLIL 250
 Qy 181 GAISYDVLAVLCPKGPLRMLVETAQERNEIFPALLIYSSAMVTVGMKLDPPSSQAGAL- 240
 Db 251 AVISYDVLAVLCPKGPLRMLVETAQERNEIFPALLIYSSAMVTVGMKLDPPSSQAGAL- 310
 Qy 240 -QLPYDPE-NEEDSYDSFGE---PSYPVEFPEPLTCYPG-----EEL----- 277
 Db 311 KNSKYNAESTERESQDTVAENDGGFSEWEAQRDHLGPHRSTPSRAAVQELSSSILA 370
 Qy 277 -EEEEERGKVLGLGDFIFYSVLVGKAAATGSGDWNNTTACFVAILIGLCLTLILLAVFKK 335
 Db 371 GEDPEERGKVLGLGDFIFYSVLVGKASATASGDWNNTTACFVAILIGLCLTLILLAVFKK 430
 Qy 336 ALPALPISITFGLIFYFSTDNLRPFMDTLASHQLYI 372
 Db 431 ALPALPISITFGLIFYFATDYLVPQMDQLAFHQFYI 467

RESULT 15
 W28507
 ID W28507 standard; Protein; 407 AA.
 AC W28507;
 DT 07-DEC-1997 (first entry)
 DE Partial AD3 sequence.
 KW AD3; AD4/AD3LP; Alzheimer's disease; chromosome; missegregation;
 KW presenilin; inhibitor; AD; trisomy 21.
 OS Homo sapiens.

FT Key Location/Qualifiers
 FT misc_difference 86
 FT /label= mutation
 FT /note= "M -> L"
 FT misc_difference 103
 FT /label= mutation
 FT /note= "H -> R"
 FT misc_difference 186
 FT /label= mutation
 FT /note= "A -> E"
 FT misc_difference 226
 FT /label= mutation
 FT /note= "L -> V"
 FT misc_difference 350

FT /label= mutation
 FT /note= "C -> Y"
 PN W09707213-A2.
 PD 27-FEB-1997.
 PF 15-AUG-1996; U13314.
 PR 16-AUG-1995; US-002448.
 PA (HARD) HARVARD COLLEGE.
 PI Li J, Potter H;
 DR WPI; 97-165297/15.
 DR N-PSDB; T87402.
 PT Identifying genes which cause chromosome missegregation - useful for
 PT identifying causes of and treatments for diseases, e.g. Alzheimer's
 PT disease, cancer and ageing
 PS Disclosure; Fig 1; 77pp; English.
 CC Identifying genes which cause improper chromosome segregation,
 CC screening for inhibitors of chromosome missegregation and processes
 CC caused by genes encoding chromosome missegregation promoters
 CC was exemplified using Alzheimer's disease. The sequences
 CC given in T87401 to T87426 can be used in the above methods.
 CC The five mutations indicated in the Features Table cosegregate
 CC with early-onset familial Alzheimer's disease. It is predicted
 CC that these mutations result in increased levels of cells with
 CC trisomy 21 in carriers of the mutation compared with non-carriers.
 SQ Sequence 407 AA;

Query Match 74.3%; Score 1429.5; DB 1; Length 407;
 Best Local Similarity 72.3%; Pred. No. 5.4e-147;
 Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

Qy 1 EELTKYGAHVIMLFVPTLCMVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRLL 60
 Db 11 EELTKYGAHVIMLFVPTLCMVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRLL 70
 Qy 61 NSVLNTLMISVIIVMTIFLVLYKYRCYKFTGHWLIMSSLMLEFLTYIYLGEVLKTYN 120
 Db 71 HSLNAAIMISVIIVMTIFLVLYKYRCYKFTGHWLIMSSLMLEFLTYIYLGEVLKTYN 130
 Qy 121 VANDYITVALLIWNFGVGMISIHKGLPLRQQAAYLIMISALMALVFIKYLPEWTAWLIL 180
 Db 131 VANDYITVALLIWNFGVGMISIHKGLPLRQQAAYLIMISALMALVFIKYLPEWTAWLIL 190
 Qy 181 GAISYDVLAVLCPKGPLRMLVETAQERNEIFPALLIYSSAMVTVGMKLDPPSSQAGAL- 240
 Db 191 AVISYDVLAVLCPKGPLRMLVETAQERNEIFPALLIYSSAMVTVGMKLDPPSSQAGAL- 250
 Qy 240 -QLPYDPE-NEEDSYDSFGE---PSYPVEFPEPLTCYPG-----EEL----- 277
 Db 251 KNSKYNAESTERESQDTVAENDGGFSEWEAQRDHLGPHRSTPSRAAVQELSSSILA 310
 Qy 277 -EEEEERGKVLGLGDFIFYSVLVGKAAATGSGDWNNTTACFVAILIGLCLTLILLAVFKK 335
 Db 311 GEDPEERGKVLGLGDFIFYSVLVGKASATASGDWNNTTACFVAILIGLCLTLILLAVFKK 370
 Qy 336 ALPALPISITFGLIFYFSTDNLRPFMDTLASHQLYI 372
 Db 371 ALPALPISITFGLIFYFATDYLVPQMDQLAFHQFYI 407

Search completed: March 20, 2000, 05:31:21
 Job time: 4210 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 19:55:31 ; Search time 26.47 seconds
(without alignments)
186.879 Million cell updates/sec

Title: US-08-509-359B-138

Perfect score: 1923

Sequence: 1 EELTLKYGAKHVMFLFVPT.....STDNLVRPFMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 133990 seqs, 13297546 residues

Database : Issued_Patents_AA.*

Word size : 0

Number of hits that pass the threshold : 133990

- 1: /cgn2_6/ptodata/2/iaa/5A_COMB.pep.*
- 2: /cgn2_6/ptodata/2/iaa/5B_COMB.pep.*
- 3: /cgn2_6/ptodata/2/iaa/6_COMB.pep.*
- 4: /cgn2_6/ptodata/2/iaa/PCTUS9_COMB.pep.*
- 5: /cgn2_6/ptodata/2/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1923	100.0	448	2	US-08-967-101-137
2	1923	100.0	372	2	US-08-967-101-138
3	1923	100.0	448	2	US-08-592-541-137
4	1923	100.0	372	2	US-08-592-541-138
5	1907.5	99.2	447	2	US-08-875-972-29
6	1591.5	82.8	376	2	US-08-875-972-2
7	1429.5	74.3	407	2	US-08-967-101-134
8	1429.5	74.3	407	2	US-08-875-972-4
9	1429.5	74.3	467	2	US-08-592-541-134
10	1429.5	74.3	467	3	US-08-670-964-2
11	1429.5	74.3	463	3	US-08-670-964-4
12	1423.5	74.0	467	2	US-08-967-101-2
13	1423.5	74.0	467	2	US-08-592-541-2
14	1412.5	73.5	463	2	US-08-670-479-18
15	1381.5	71.8	467	2	US-08-967-101-4
16	1381.5	71.8	467	2	US-08-592-541-4
17	1127.5	58.6	541	2	US-08-967-101-166
18	1127.5	58.6	541	2	US-08-592-541-166
19	100	5.2	1294	2	US-08-819-288-3
20	98	5.1	1321	1	US-08-261-822A-3
21	98	5.1	1321	4	PCT-US95-07744A-3
22	97	5.0	1334	2	US-08-996-545-2
23	89	4.6	509	2	US-09-031-392-6
24	88.5	4.6	452	1	US-08-117-361C-1
25	88.5	4.6	3169	2	US-08-477-451-6
26	87	4.5	492	2	US-08-355-844-3
27	87	4.5	492	4	PCT-US95-16126-3
28	82	4.3	502	1	US-08-484-840-3
29	82	4.3	1873	1	US-08-336-257A-7
30	82	4.3	502	1	US-08-483-094-3
31	82	4.3	413	2	US-08-808-793-25
32	81	4.2	2100	2	US-08-808-793-23
33	80	4.2	837	1	US-07-923-976-2
34	80	4.2	602	1	US-08-295-814E-2

ALIGNMENTS

RESULT 1

US-08-967-101-137

; Sequence 137, Application US/08967101

; Patent No. 5840340

; GENERAL INFORMATION:

; APPLICANT: ST. GEORGE-HYSLOP, PETER H

; APPLICANT: ROMMENS, JOHANNA M

; APPLICANT: FRASER, PAUL E

; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED

; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE

; NUMBER OF SEQUENCES: 183

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: TESTA, HURWITZ & THIBEAULT

; STREET: High Street Tower - 125 High Street

; CITY: Boston

; STATE: Massachusetts

; COUNTRY: U.S.A.

; ZIP: 02110

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent in Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; FILING DATE: 10-NOV-1997

; CLASSIFICATION: 435

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/592,541

; FILING DATE:

; ATTORNEY/AGENT INFORMATION:

; NAME: Pitcher, Edmund R.

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (617) 248-7000

; TELEFAX: (617) 248-7100

; INFORMATION FOR SEQ ID NO: 137:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 448 amino acids

; TYPE: amino acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: protein

; US-08-967-101-137

Query Match 100.0%; Score 1923; DB 2; Length 448;

Best Local Similarity 100.0%; Pred. No. 2.2e-190;

Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 EELTLKYGAKHVMFLFVPTLCMIVVATIKSVRFYTERNGQLIYPTFTDPSVGQRL 60

Db 77 EELTLKYGAKHVMFLFVPTLCMIVVATIKSVRFYTERNGQLIYPTFTDPSVGQRL 136

QY 61 NSVLATLIMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 120

Db 137 NSVLATLIMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 196

Db 77 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFYTEKNGQLIYTPFTEDPSVGQRL 136
Qy 61 NSVLTLMISVIVVMTIFLVVLYKYRCYKFIHGWLMISLMMLLFYIYLGEVLKTYN 120
Db 137 NSVLTLMISVIVVMTIFLVVLYKYRCYKFIHGWLMISLMMLLFYIYLGEVLKTYN 196
Qy 121 VMDYPTLLTWNFGAVGVCVCIHMKGPLVLQQAAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 197 VMDYPTLLTWNFGAVGVCVCIHMKGPLVLQQAAYLIMISALMALVFIKYLPEWSAWVIL 256
Qy 181 GAISYVDLVAVLCPKGPLRMLVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQGALQ 240
Db 257 GAISYVDLVAVLCPKGPLRMLVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQGALQ 316
Qy 241 LPYDPEMEDSDSDSGEPSPYVFPPLTGYGPEEEERGVKLGDFIFYSVLVGK 300
Db 317 LPYDPEMEDSDSDSGEPSPYVFPPLTGYGPEEEERGVKLGDFIFYSVLVGK 376
Qy 301 AAATGSGDWNNTLACFVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
Db 377 AAATGSGDWNNTLACFVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 436
Qy 361 FMDTLASHOLYI 372
Db 437 FMDTLASHOLYI 448

RESULT 4
US-08-592-541-138
; Sequence 138, Application US/08592541
; Patent No. 5986054
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 138:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 372 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-592-541-138

Query Match 100.0%; Score 1923; DB 2; Length 372;
Best Local Similarity 100.0%; Pred. No. 1.7e-190;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFYTEKNGQLIYTPFTEDPSVGQRL 60
Db 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFYTEKNGQLIYTPFTEDPSVGQRL 60
Qy 61 NSVLTLMISVIVVMTIFLVVLYKYRCYKFIHGWLMISLMMLLFYIYLGEVLKTYN 120
Db 61 NSVLTLMISVIVVMTIFLVVLYKYRCYKFIHGWLMISLMMLLFYIYLGEVLKTYN 120
Qy 121 VMDYPTLLTWNFGAVGVCVCIHMKGPLVLQQAAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 121 VMDYPTLLTWNFGAVGVCVCIHMKGPLVLQQAAYLIMISALMALVFIKYLPEWSAWVIL 180
Qy 181 GAISYVDLVAVLCPKGPLRMLVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQGALQ 240
Db 181 GAISYVDLVAVLCPKGPLRMLVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQGALQ 240
Qy 241 LPYDPEMEDSDSDSGEPSPYVFPPLTGYGPEEEERGVKLGDFIFYSVLVGK 300
Db 241 LPYDPEMEDSDSDSGEPSPYVFPPLTGYGPEEEERGVKLGDFIFYSVLVGK 300
Qy 301 AAATGSGDWNNTLACFVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
Db 301 AAATGSGDWNNTLACFVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
Qy 361 FMDTLASHOLYI 372
Db 361 FMDTLASHOLYI 372

RESULT 5
US-08-875-972-29
; Sequence 29, Application US/08875972
; Patent No. 5985564
; GENERAL INFORMATION:
; APPLICANT: Huntington Potter and Jinhue Li
; TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
; TITLE OF INVENTION: CHROMOSOME NON-DISJUNCTION
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02173-4799
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/875,972
; FILING DATE: 08-AUG-97
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/002,448
; FILING DATE: 16-AUG-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan Esq., Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: HU95-03PA
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781) 861-6240
; TELEFAX: (781) 861-9540
; INFORMATION FOR SEQ ID NO: 29:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 447 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-875-972-29

TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-967-101-134

Query Match 74.3%; Score 1429.5; DB 2; Length 467;
Best Local Similarity 72.3%; Pred. No. 2e-139;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

QY 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYTPPTDTPSVGQRLL 60
DB 71 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYTPPTDTPSVGQRAL 130
QY 61 NSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLMSSLMFLFYIYLGEVLKTYN 120
DB 131 HSLNAAIMISIVVMTILLVLYKYRCYKVIHAWLIISLILFFFIYIYLGEVFKTYN 190
QY 121 VAMDYPTLLTWNFGAVGMCVHMKGPLVLOQAYLIMISALMALVFYKYLPEWSAWIL 180
DB 191 VAVDIITVALLIWNFGVGMISIHMKGPLRLOQAYLIMISALMALVFYKYLPEWTAWIL 250
QY 181 GAISYDLVAVLCPLKPLMLVETAQERNEIFFPALIISSANVTVGMKLDPSQOGAL- 240
DB 251 AVISYDLVAVLCPLKPLMLVETAQERNEIFFPALIISSANVTVGMKLDPSQOGAL- 310
QY 240 -QLPYDPE-MEEDSYDSFGE---PSYPEVFEPPLTGYPG-----EEL----- 277
DB 311 KNSKYNAESTERESQDTVAENDDGGFSEWEAQRDHSHLPHRSTPESRAAVQELSSILA 370
QY 277 -EEEEERGVKLGDFIFYSVLVGRKAAATGSGDNTTACFVAILIGLCLTLLLAFAFKK 335
DB 371 GEDPEERGVKLGDFIFYSVLVGRKAAATGSGDNTTACFVAILIGLCLTLLLAFAFKK 430
QY 336 ALPALPISITFGLIFEFSTDLNVRPMDTLASHOLYI 372
DB 431 ALPALPISITFGLVYFATDYLVPQMDQLAFHFYI 467

RESULT 8

US-08-875-972-4
Sequence 4, Application US/08875972
Patent No. 5985564

GENERAL INFORMATION:
APPLICANT: Huntington Potter and Jinhue Li
TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
TITLE OF INVENTION: CHROMOSOME NON-DISJUNCTION
NUMBER OF SEQUENCES: 29
CORRESPONDENCE ADDRESS:
ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
STREET: Two Militia Drive
CITY: Lexington
STATE: Massachusetts
COUNTRY: USA
ZIP: 02173-4799

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
APPLICATION NUMBER: US/08/875,972
FILING DATE: 08-AUG-97
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/002,448
FILING DATE: 16-AUG-1995
ATTORNEY/AGENT INFORMATION:
NAME: Granahan Esq., Patricia
REGISTRATION NUMBER: 32,227
REFERENCE/DOCKET NUMBER: H095-03PA
TELECOMMUNICATION INFORMATION:

TELEPHONE: (781) 861-6240
TELEFAX: (781) 861-9540
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 407 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-875-972-4

Query Match 74.3%; Score 1429.5; DB 2; Length 407;
Best Local Similarity 72.3%; Pred. No. 1.6e-139;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

QY 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYTPPTDTPSVGQRLL 60
DB 11 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYTPPTDTPSVGQRAL 70
QY 61 NSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLMSSLMFLFYIYLGEVLKTYN 120
DB 71 HSLNAAIMISIVVMTILLVLYKYRCYKVIHAWLIISLILFFFIYIYLGEVFKTYN 130
QY 121 VAMDYPTLLTWNFGAVGMCVHMKGPLVLOQAYLIMISALMALVFYKYLPEWSAWIL 180
DB 131 VAVDIITVALLIWNFGVGMISIHMKGPLRLOQAYLIMISALMALVFYKYLPEWTAWIL 190
QY 181 GAISYDLVAVLCPLKPLMLVETAQERNEIFFPALIISSANVTVGMKLDPSQOGAL- 240
DB 191 AVISYDLVAVLCPLKPLMLVETAQERNEIFFPALIISSANVTVGMKLDPSQOGAL- 250
QY 240 -QLPYDPE-MEEDSYDSFGE---PSYPEVFEPPLTGYPG-----EEL----- 277
DB 251 KNSKYNAESTERESQDTVAENDDGGFSEWEAQRDHSHLPHRSTPESRAAVQELSSILA 310
QY 277 -EEEEERGVKLGDFIFYSVLVGRKAAATGSGDNTTACFVAILIGLCLTLLLAFAFKK 335
DB 311 GEDPEERGVKLGDFIFYSVLVGRKAAATGSGDNTTACFVAILIGLCLTLLLAFAFKK 370
QY 336 ALPALPISITFGLIFEFSTDLNVRPMDTLASHOLYI 372
DB 371 ALPALPISITFGLVYFATDYLVPQMDQLAFHFYI 407

RESULT 9

US-08-592-541-134
Sequence 134, Application US/08592541
Patent No. 5986054

GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183
CORRESPONDENCE ADDRESS:
ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/592,541
FILING DATE:
CLASSIFICATION: 800
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.

OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/670,964
FILING DATE: 26-JUN-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/001,142
FILING DATE: 13-JUL-1995
APPLICATION NUMBER: 60/001,501
FILING DATE: 18-JUL-1995
ATTORNEY/AGENT INFORMATION:
NAME: Han, William T
REGISTRATION NUMBER: 34,344
REFERENCE/DOCKET NUMBER: P50358
TELECOMMUNICATION INFORMATION:
TELEPHONE: 610-270-5219
TELEFAX: 610-270-5090
TELEX:
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 463 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-670-964-4

Query Match 74.3%; Score 1429.5; DB 3; Length 463;
Best Local Similarity 72.3%; Pred. No. 1.9e-139;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

QY 1 BELTLKYGAKHVMFLFVPVTLQAVYIMISALMALVFIKYLPEWSAWVIL 60
DB 67 BELTLKYGAKHVMFLFVPVTLQAVYIMISALMALVFIKYLPEWSAWVIL 126
QY 61 NSVLTNLMISIVVMTFLVLYKYRCYKFIHGLWIMSLMLFLFYIYLGEVLKYN 120
DB 127 HSLNLAAMISIVVMTILLVLYKYRCYKFIHAWLIISLSSLLFFSFYILGEVFKYN 186
QY 121 VAMDYPTLLTWNFGAVGMVCIHMKGPLVLOQAVYIMISALMALVFIKYLPEWSAWVIL 180
DB 187 VAVDYITVALLIWNFGVGMISIHMKGPLRLOQAVYIMISALMALVFIKYLPEWTAWLIL 246
QY 181 GAISYDVLVAVLCPRGLRMLVETAQERNEPIFPALIISSAMVWTVMGAKLPSSQGAL- 240
DB 247 AVISYDVLVAVLCPRGLRMLVETAQERNEPIFPALIISSAMVWTVMGAKLPSSQGAL- 306
QY 240 -QLPDPE-MEEDSDSFGE---PSYPEVFEPPLTGYPG-----EEL----- 277
DB 307 KNSKYNAESTERESODTVAENDDDGGFSEWEAQRDHLGPHRSTPESRAAVQELSSILA 366
QY 277 -EEERGVKLGIDFIFYSVLVGKAAATGSDWNTTACFVAILIGLCTLLLLAVFKK 335
DB 367 GEDPEERGKLGIDFIFYSVLVGKASATASGDWNTTACFVAILIGLCTLLLLAIFKK 426
QY 336 ALPALPISITFGLIFVFTDNLVRPMDTLASHOLYI 372
DB 427 ALPALPISITFGLVYFATDYLQVPMQDLAFHQFYI 463

RESULT 12
US-08-967-101-2
Sequence 2, Application US/08967101
Patent No. 5840540
GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:
ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,101
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Fitcher, Edmund R.
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 467 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-967-101-2

Query Match 74.0%; Score 1423.5; DB 2; Length 467;
Best Local Similarity 72.0%; Pred. No. 8.2e-139;
Matches 286; Conservative 33; Mismatches 53; Indels 25; Gaps 5;

QY 1 BELTLKYGAKHVMFLFVPVTLQAVYIMISALMALVFIKYLPEWSAWVIL 60
DB 71 BELTLKYGAKHVMFLFVPVTLQAVYIMISALMALVFIKYLPEWSAWVIL 130
QY 61 NSVLTNLMISIVVMTFLVLYKYRCYKFIHGLWIMSLMLFLFYIYLGEVLKYN 120
DB 131 HSLNLAAMISIVVMTILLVLYKYRCYKFIHAWLIISLSSLLFFSFYILGEVFKYN 190
QY 121 VAMDYPTLLTWNFGAVGMVCIHMKGPLVLOQAVYIMISALMALVFIKYLPEWSAWVIL 180
DB 191 VAVDYITVALLIWNFGVGMISIHMKGPLRLOQAVYIMISALMALVFIKYLPEWTAWLIL 250
QY 181 GAISYDVLVAVLCPRGLRMLVETAQERNEPIFPALIISSAMVWTVMGAKLPSSQGAL- 240
DB 251 AVISYDVLVAVLCPRGLRMLVETAQERNEPIFPALIISSAMVWTVMGAKLPSSQGAL- 310
QY 240 -QLPDPE-MEEDSDSFGE---PSYPEVFEPPLTGYPG-----EEL----- 277
DB 311 KNSKYNAESTERESODTVAENDDDGGFSEWEAQRDHLGPHRSTPESRAAVQELSSILA 370
QY 277 -EEERGVKLGIDFIFYSVLVGKAAATGSDWNTTACFVAILIGLCTLLLLAVFKK 335
DB 371 GEDPEERGKLGIDFIFYSVLVGKASATASGDWNTTACFVAILIGLCTLLLLAIFKK 430
QY 336 ALPALPISITFGLIFVFTDNLVRPMDTLASHOLYI 372
DB 431 ALPALPISITFGLVYFATDYLQVPMQDLAFHQFYI 467

RESULT 13
US-08-592-541-2
Sequence 2, Application US/08592541
Patent No. 5986054
GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H

Db 367 GEDPEERGKVLGLDFIFYSVLVGRASATASGDWNTTIACFVAILIGLCLTLLLLAIFKK 426
Qy 336 ALPALPISITGLFIFYSTDNIVRFMDTFLASHQIYI 372
Db 427 ALPALPISITGLFVFATDYLVQPFMDQLAFHQFYI 463
RESULT 15
US-08-967-101-4
; Sequence 4, Application US/08967101
; Patent No. 5840540
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/967,101
; FILING DATE: 10-NOV-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/592,541
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 467 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-967-101-4

Query Match 71.8%; Score 1381.5; DB 2; Length 467;
Best Local Similarity 68.7%; Pred. No. 1.8e-134;
Matches 274; Conservative 38; Mismatches 58; Indels 29; Gaps 4;
Qy 1 BELTLKYGAKHIVLFPVTLICMIVVATIKSVRYTEKNGOLIYTPETDTPSVQRL 60
Db 71 BELTLKYGAKHIVLFPVTLICMIVVATIKSVRYTEKNGOLIYTPETDTPSVQRL 130
Qy 61 NSVLNTLMISIVVMITFLVLYKYRCYKFIHGLIMSSMLLFLFYIYLGVLKTYN 120
Db 131 HSILNAAIMISIVIMTLLVLYKYRCYKVIHAWLIISLLEFFSFIVLGEVFKTYN 190
Qy 121 VMDYPTLLLVNFGVAGMCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 180
Db 191 VXDVTVTALLIWNWVGVIHAWKGPLRLOQAYLIMISALMALVFIKYLPEWTAWIL 250
Qy 181 GAISVYDLVAVLCPKGPLMVLVETAQERNETLFPALIIYSSAMVTVGMKLDPPSSOGALQ 240
Db 251 AVISYVDLVAVLCPKGPLMVLVETAQERNETLFPALIIYSSAMVTVGMKLDPPSSOGALQ 308

Qy 241 LPYDPEME-----EDSYDSFGPSYVEPEPPLTGYPG-----EEL----- 277
Db 309 YPKNPKYNTQRAERETQDSGSGNDGGFSEWEAQDSDHLGPHRSTPESRAAVQELSGSI 368
Qy 277 ---EEEEERGVKGLGDFIFYSVLVGRASATASGDWNTTIACFVAILIGLCLTLLLLAVF 333
Db 369 LTSDEPPEERGKVLGLGDFIFYSVLVGRASATASGDWNTTIACFVAILIGLCLTLLLLAIY 428
Qy 334 KKALPALPISITGLFIFYSTDNIVRFMDTFLASHQIYI 372
Db 429 KKGXPAXPISITGLFVFATDYLVQPFMDQLAFHQFYI 467

Search completed: March 18, 2000, 19:55:32
Job time: 3230 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 14:11:52 ; Search time 41.25 Seconds
(without alignments)
425.381 Million cell updates/sec

Title: US-08-509-359B-138
Perfect score: 1923
Sequence: 1 EELTKYGAHVIMLFVPT.....STDNLVRPFMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 142080 seqs, 47169319 residues

Database : PIR_62.*

Word size : 0

Number of hits that pass the threshold : 142080

- 1: pir1.*
- 2: pir2.*
- 3: pir3.*
- 4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	1923	100.0	448	2	I58098	ES-1 protein - hum
2	1915	99.6	448	2	A56993	presenilin 2 - hum
3	1802.5	93.7	442	2	J39174	seven trans-membra
4	1534	79.8	449	2	J5391	presenilin-beta -
5	1431.5	74.4	467	2	I78388	S182 protein - mou
6	1431.5	74.4	433	2	JC5390	presenilin-alpha -
7	1429.5	74.3	467	2	S58396	presenilin 1, spli
8	1429.5	74.3	463	2	S63683	presenilin I-463 -
9	1413.5	73.5	463	2	JC5081	presenilin 1 prote
10	1413.5	73.5	467	2	JC5080	presenilin 1 prote
11	1011	52.6	374	2	S63684	presenilin 1, spli
12	956	49.7	461	2	S60253	sel-12 protein - C
13	524.5	27.3	358	2	T15184	hypothetical prote
14	513	26.7	453	2	T00724	presenilin homolog
15	274	14.2	465	2	A43459	sperm membrane pro
16	110	5.7	2016	2	A38195	sodium channel pro
17	104	5.4	826	2	T02268	potassium transpor
18	104	5.4	398	2	H75043	mg2+ transport pro
19	103	5.4	382	2	S47882	ubiquinol--cytochr
20	102	5.3	379	2	I48135	ubiquinol--cytochr
21	101.5	5.3	1681	2	A55138	sodium channel mna
22	100.5	5.2	324	2	S36646	integrin-associate
23	99.5	5.2	1840	1	CHRTM1	sodium channel pro
24	99.5	5.2	379	2	JC6178	serotonin receptor
25	99	5.1	2019	2	A33996	sodium channel pro
26	98.5	5.1	531	2	T11596	hypothetical prote
27	98	5.1	461	2	T11829	NADH dehydrogenase
28	98	5.1	217	2	S01095	hypothetical prote
29	97	5.0	381	2	T11440	ubiquinol--cytochr
30	97	5.0	447	2	S52968	NADH dehydrogenase
31	97	5.0	238	2	S02063	H+-transporting AT
32	97	5.0	441	2	S13425	endothelin recepto
33	97	5.0	299	2	D65187	ubiquinol--cytochr
34	96.5	5.0	308	2	S22928	rad3 protein - fis
35	96.5	5.0	1070	2	S25834	

36 96 5.0 507 2 B64433 probable O-antigen
37 96 5.0 592 2 E70488 cytochrome-c oxida
38 95.5 5.0 488 1 QXASM4 NADH dehydrogenase
39 95.5 5.0 442 1 JQ1042 endothelin recepto
40 95.5 5.0 294 2 D53290 oligopeptide trans
41 95.5 5.0 420 2 A47649 probable inner mem
42 95.5 5.0 768 2 S52684 probable membrane
43 95.5 5.0 799 2 H71255 probable cell divi
44 95 4.9 492 2 A30797 glucose transport
45 95 4.9 484 2 S75022 bacteriochlorophyl

ALIGNMENTS

RESULT 1
I58098
E5-1 protein - human
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 29-Sep-1999
C:Accession: I58098
R:Rogaev, E.I.; Sherrington, R.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Liang, Y.; C
.; Cohen, D.; Lannfelt, L.; Fraser, P.E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 376, 775-778, 1995
A:Title: Familial Alzheimer's disease in kindreds with missense mutations in a gene o
A:Reference number: I58098; MUID:95379971
A:Accession: I58098
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: mRNA
A:Residues: 1-448 <RES>
A:Cross-references: GB:L44577; NID:g950347; PIDN:AAC42012.1; PID:g950348
C:Genetics:
A:Gene: E5-1
C:Superfamily: presenilin

Query Match 100.0%; Score 1923; DB 2; Length 448;
Best Local Similarity 100.0%; Pred. No. 3.5e-138;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLIYTPFTEDTPSVGQRL 60
DB 77 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLIYTPFTEDTPSVGQRL 136
QY 61 NSVLTLMISVIVVMTIFLVVLYKYRCYKFTHGWLINSSLMFLFYIYLGEVLKTYN 120
DB 137 NSVLTLMISVIVVMTIFLVVLYKYRCYKFTHGWLINSSLMFLFYIYLGEVLKTYN 196
QY 121 VMDYPTLLLTWNFGVGMVCIHWKGPLVLOQAAYLIMISALMALVFIKYLPEWSAWVIL 180
DB 197 VMDYPTLLLTWNFGVGMVCIHWKGPLVLOQAAYLIMISALMALVFIKYLPEWSAWVIL 256
QY 181 GAISVYDLAVLCPKPLRMVETAQERNEFPFALITYSSAMVTVGMKLPSSOGALQ 240
DB 257 GAISVYDLAVLCPKPLRMVETAQERNEFPFALITYSSAMVTVGMKLPSSOGALQ 316
QY 241 LPYPMEEDSYDSGEPSEYFEPPLTGYGPEEEEEEERGVKLGDFIFYSVLVGK 300
DB 317 LPYPMEEDSYDSGEPSEYFEPPLTGYGPEEEEEEERGVKLGDFIFYSVLVGK 376
QY 301 AAATGSGDWTTLACFAVILGLCTLALLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
DB 377 AAATGSGDWTTLACFAVILGLCTLALLAVFKKALPALPISITFGLIFYFSTDNLVRP 436
QY 361 FMDTLASHQLYI 372
DB 437 FMDTLASHQLYI 448

RESULT 2
A56993
presenilin 2 - human
N:Alternate names: Alzheimer's disease protein 4

Qy 240 -OLPY-----DPEMEDSYDSFGPEPSYVEFPPTLTGYPGELEEEERGVKLGIDRFY 294
Db 319 QOVQHIDRNPTEGANSIVDEAAETRIQ-----TQSLSSDDPEERGVKLGIDRFY 371
Qy 295 SVLVGKAAATGSGDWNNTLACFVAILIGLCITLLALLAVFKKALPALPISITIFGLIFEST 354
Db 372 SVLVGKAAATGSGDWNNTLACFVAILIGLCITLLALLAVFKKALPALPISITIFGLIFEST 431
Qy 355 DNLVRPMDTLASHQLYI 372
Db 432 DNLVRPMDTLASHQMYI 449
RESULT 5
I78388
S182 protein - mouse
C:Species: Mus musculus (house mouse)
C:Date: 27-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 29-Sep-1999
C:Accession: I78388
R:Sherrington, R.; Rogaev, E.I.; Liang, Y.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Chl.
ero, I.; Pinessi, L.; Nee, L.; Chumakov, I.; Pollen, D.; Brookes, A.; Sanseau, P.; Polin
E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 375, 754-760, 1995
A:Title: Cloning of a gene bearing missense mutations in early-onset familial Alzheimer
A:Reference number: I58095; MUID:95319502
A:Accession: I78388
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-467 <RES>
A:Cross-references: GB:L42177; NID:g904129; PIDN:AAC42094.1; PID:g904130
C:Superfamily: presenilin
Query Match 74.4%; Score 1431.5; DB 2; Length 467;
Best Local Similarity 71.2%; Pred. No. 5.1e-101;
Matches 284; Conservative 36; Mismatches 50; Indels 29; Gaps 4;
Qy 1 EELTLKYGAKHVMFLFVPVTLICMIVVATIKSVRYTEKNGQLIYTPPTEDTPSVGQRLL 60
Db 71 EELTLKYGAKHVMFLFVPVTLICMIVVATIKSVSFYTRKDGQLIYTPPTEDTETVGQRAL 130
Qy 61 NSVLNTLMISVIVMTIFVLVLYKYCYKFIHGWLMISLMLFLFYIYLGEVLKTYN 120
Db 131 HSLUNAIMSVIVMTILLVLYKYCYKVIHAWLISSLLLLFFSFYILGEVFKTYN 190
Qy 121 VAMDYPTLLLTVMNFGAVGVCIHKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 191 VANDYPTVALLIWNFGVGMIAIHWKGPLRLOQAYLIMISALMALVFIKYLPEWTAWLIL 250
Qy 181 GAISVYDLVAVLCPKPLMLVETAQERNEPFPALIIYSAMVTVGMKALDPS 240
Db 251 AVISYDLVAVLCPKPLMLVETAQERNEPFPALIIYSSTMTWLVNMAEGDPEAQ--RR 308
Qy 241 LPYDPENE-----EDSYDSFGPEPSYVEFPPLTGYPG-----EEL----- 277
Db 309 VPKNPKYNTQARERTQDSGSGNDGDFSEWEAQRDHSLGPHRSTPESRAVQELSGSI 368
Qy 277 ---EEERGVKLGIDFIFYSVLVGKAAATGSGDWNNTLACFVAILIGLCITLLALLAVF 333
Db 369 LTSEDPERGVKLGIDFIFYSVLVGKASATASGDWNTTACFVAILIGLCITLLALLAIF 428
Qy 334 KKALPALPISITIFGLIFFYFSTDNLVRPMDTLASHQLYI 372
Db 429 KKALPALPISITIFGLVFFATDYLVPQFMDQLAFHQFYI 467
RESULT 6
JC5390
presenilin-alpha - African clawed frog
C:Species: Xenopus laevis (African clawed frog)
C:Date: 04-Jun-1997 #sequence_revision 18-Jul-1997 #text_change 29-Sep-1999
C:Accession: JC5390
R:Tsujiura, A.; Yasojima, K.; Hashimoto-Gotoh, T.

Biochem. Biophys. Res. Commun. 231, 392-396, 1997
A:Title: Cloning of Xenopus presenilin-alpha and -beta cDNAs and their differential e
A:Reference number: JC5390; MUID:97223465
A:Accession: JC5390
A>Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-433 <TSU>
A:Cross-references: DBJ:D84427; NID:gl944353; PIDN:BAAL9570.1; PID:dl020347; PID:gl9
A:Experimental source: brain
C:Comment: This protein plays a role in negative regulation of apoptotic cascades dur
C:Superfamily: presenilin
F:48-66/Domain: transmembrane #status predicted <TM1>
F:99-119/Domain: transmembrane #status predicted <TM2>
F:130-149/Domain: transmembrane #status predicted <TM3>
F:161-178/Domain: transmembrane #status predicted <TM4>
F:187-203/Domain: transmembrane #status predicted <TM5>
F:210-227/Domain: transmembrane #status predicted <TM6>
F:374-394/Domain: transmembrane #status predicted <TM7>
Query Match 74.4%; Score 1431.5; DB 2; Length 433;
Best Local Similarity 72.0%; Pred. No. 4.7e-101;
Matches 286; Conservative 27; Mismatches 59; Indels 25; Gaps 4;
Qy 1 EELTLKYGAKHVMFLFVPVTLICMIVVATIKSVRYTEKNGQLIYTPPTEDTPSVGQRLL 60
Db 37 EELTLKYGAKHVMFLFVPVTLICMIVVATIKSVSFYTRFDGQLIYTPPTEDTESVQGRAL 96
Qy 61 NSVLNTLMISVIVMTIFVLVLYKYCYKFIHGWLMISLMLFLFYIYLGEVLKTYN 120
Db 97 NSILNATMISVIVMTILLVLYKYCYKVIHGWLMISLMLFLFYIYLGEVFKTYN 156
Qy 121 VAMDYPTLLLTVMNFGAVGVCIHKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 157 VANDYPTTALLIWNFGVGMICIHKGPLLOQAYLIMISALMALVFIKYLPEWTWLLIL 216
Qy 181 GAISVYDLVAVLCPKPLMLVETAQERNEPFPALIIYSAMVTVGMKALDPS-SQGL 239
Db 217 AVISYDLVAVLSPKPLMLVETAQERNEPFPALIIYSSTMTWLVNMAADGDLKQAS 276
Qy 240 QLPYDPEN-----EDSYDSFGPEPSYVEFP-----PLTGYPGCEL----- 277
Db 277 TKTYTQAPTAPHRSDSAASDNDGDFDTTWDHNAQIGPINSTPESRVAQALPNSPP 336
Qy 277 ---EEERGVKLGIDFIFYSVLVGKAAATGSGDWNNTLACFVAILIGLCITLLALLAVFK 335
Db 337 SEDPERGVKLGIDFIFYSVLVGKASATASGDWNTTACFVAILIGLCITLLALLAIFK 396
Qy 336 ALPALPISITIFGLIFFYFSTDNLVRPMDTLASHQLYI 372
Db 397 ALPALPISITIFGLVFFATDYLVPQFMDQLAFHQFYI 433
RESULT 7
S58396
presenilin 1, splice form 467 - human
N:Alternate names: Alzheimer's disease protein 3; protein s182
C:Species: Homo sapiens (man)
C:Date: 29-Jan-1998 #sequence_revision 13-Feb-1998 #text_change 29-Sep-1999
C:Accession: S58396; S71401; S71402
R:Sherrington, R.; Rogaev, E.I.; Liang, Y.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; C
ero, I.; Pinessi, L.; Nee, L.; Chumakov, I.; Pollen, D.; Brookes, A.; Sanseau, P.; Po
E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 375, 754-760, 1995
A:Title: Cloning of a gene bearing missense mutations in early-onset familial Alzheim
A:Reference number: I58095; MUID:95319502
A:Accession: S58396
A:Molecule type: mRNA
A:Residues: 1-467 <SHE>
A:Cross-references: EMBL:L42110; NID:g904118; PIDN:AAB46416.1; PID:g904119
R:Vidal, R.; Ghiso, J.; Wisniewski, T.; Frangione, B.
FEBS Lett. 393, 19-23, 1996
A:Title: Alzheimer's presenilin 1 gene expression in platelets and megakaryocytes. Id

|||||
Db 369 KLGGLGDEIFYSVLGRAAMY---DLMTVYACYLAISGLGCTLLILSVYNRALPALPISI 425
QY 345 TFGILFYFSTDNLVRPFM 362
Db 426 MLGVVVFYFLTRLLMEPFV 443
RESULT 15
A43459
sperm membrane protein spe-4 - Caenorhabditis elegans
N:Alternate names: probable integral membrane protein
C:Species: Caenorhabditis elegans
C>Date: 10-Jun-1993 #sequence.Revision 18-Nov-1994 #text_change 09-Sep-1997
C:Accession: A43459; S24632; S24633
R:L'Hernault, S.W.; Arduengo, P.M.
J. Cell Biol. 119, 55-68, 1992
A:Title: Mutation of a putative sperm membrane protein in Caenorhabditis elegans prevents
A:Reference number: A43459; MUID:92407040
A:Accession: A43459
A>Status: preliminary; not compared with conceptual translation
A:Molecule type: DNA; mRNA
A:Residues: 1-465 <LHE>
A:Cross-references: EMBL:Z14067; NID:g6868; PID:g6869; EMBL:Z14066; NID:g6870; PID:g6871
A:Experimental source: strain Bristol N2
A:Note: the nucleotide sequence was submitted to the EMBL Data Library, July 1992
C:Genetics:
A:Introns: 69/3; 154/3; 200/1; 224/3; 300/1; 386/1; 435/1

Query Match 14.28; Score 274; DB 2; Length 465;
Best Local Similarity 21.58; Pred. No. 1.5e-13;
Matches 93; Conservative 80; Mismatches 127; Indels 132; Gaps 14;
QY 38 EKNQGLIYPTFEDT--PSVGORLINSVLN---TLIMISIVVMVTFVLVLYKRCYKF 91
Db 42 EVNSELSKTYFLDFSFQTTGNLLLDGFGINGVGTILVGCVSFIMLAF--VLFDPR--RI 97
QY 92 IHGWLIMSSMLLF-----LFTYIYLGVLKTYNVANDYPTLLL-----TWNFGA 137
Db 98 VKAWLTLSCLLILFGVSAQTLHDMFSQVFDQDDNNQY-----YMTIVLIVVPTVYVYGF- 152
QY 138 VGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWILGALISVYDLVAVLCPKGP 197
Db 152 -GIYAFPSNSSLIHLQIFVVTNCSLISVYLRVPFSKTFVFLWVFLWDLFAVLAPMGP 210
QY 198 LRMIVETAQERNEPIFPALYSSAMVTVGMKLDPSQG-----ALQLPYDPE 246
Db 211 LKKVOEKASDYKCVNLIMFSANEKRLTAGSNEETNEGEESIRRVKOTIEYTTRE 270
QY 247 MEEDSY-----DSF-----GEPSPYEVFPPLTGYPGEELEEE--- 282
Db 271 AQDDFYQKIRQRAAINPDSVPTSEHSPLEAEPSPIELKEKNST-----EELSDDESOTS 326
QY 282 -----R 282
Db 327 ETSSGNSLSSDSTTVSTSDISTAECDQKWDLVNSLNPNNDKRPATAADALNDGE 386
QY 283 GVKLGLGDFIFYSVLVGKAAATGSDWNTTLACFVAIILGLCTLLLLAVFKKALPALPI 342
Db 387 VLRGLGDFIFYSVLLIGQAAAGCP--FAVISALGILFLGLVVLTVTFSTEESTPALPL 444
QY 343 SITEGLIFYFST 354
Db 445 PVICGTCFYFS 456

Result No.	Score	Query			DB	ID	Description
		Match	Length	Match			
1	1923	100.0	448	1	PSN2_HUMAN	P49810	homo sapien
2	1876	97.6	448	1	PSN2_MOUSE	O61144	mus musculus
3	1866	97.0	448	1	PSN2_RAT	O88777	rattus norv
4	1855	96.5	445	1	PSN2_MICMU	P79801	microcebus
5	1534	79.8	449	1	PSN2_XENLA	O12977	xenopus lae
6	1431.5	74.4	467	1	PSN1_MOUSE	P49769	mus musculus
7	1431.5	74.4	433	1	PSN1_XENLA	O12976	xenopus lae
8	1431	74.4	468	1	PSN1_RAT	P97887	rattus norv
9	1429.5	74.3	467	1	PSN1_HUMAN	P49768	homo sapien
10	1413.5	73.5	467	1	PSN1_MICMU	P79802	microcebus
11	1127.5	58.6	541	1	PSN_DROME	O02194	drosophila
12	1032	53.7	836	1	YLAK_CAEEL	Q20076	caenorhabdi
13	972	50.5	461	1	SE12_CAEEL	P52166	caenorhabdi
14	524.5	27.3	358	1	HOP1_CAEEL	O02100	caenorhabdi
15	513	26.7	453	1	PSNH_ARATH	O64668	arabidopsis
16	274	14.2	465	1	SPB4_CAEEL	Q01608	caenorhabdi
17	110	5.7	2016	1	CIN5_HUMAN	I44524	homo sapien
18	108	5.6	381	1	CYB_DASCR	Q43302	dasyercus
19	103	5.4	382	1	CYB_DIDMA	P34303	didelphis m
20	103	5.4	380	1	CYB_MICLO	P56731	microtus l
21	102.5	5.3	381	1	CYB_ANTFL	Q33706	antechinus
22	102	5.3	381	1	CYB_PASMA	O03522	dasyurus m
23	102.5	5.3	381	1	CYB_PSENI	Q35553	pseudantech
24	101.5	5.3	381	1	CYB_NINYP	Q35196	ningau i yvo
25	101.5	5.3	381	1	CYB_PARAP	Q35377	parantechin
26	101	5.3	381	1	CYB_PLAMS	Q35533	planigale m
27	100.5	5.2	381	1	CYB_PSEMD	O03543	pseudantech
28	100	5.2	381	1	CYB_DSASGE	O20604	dasyurus ge
29	99.5	5.2	1840	1	CIN4_RAT	P15390	rattus norv
30	99	5.1	2019	1	CIN5_RAT	P15389	rattus norv
31	99	5.1	381	1	CYB_ANTMI	O63534	antechinus
32	99	5.1	381	1	CYB_PHATA	Q35673	phascogale
33	98.5	5.1	531	1	YDFG_SCHPO	Q10487	schizosacch
34	98	5.1	381	1	CYB_ANTME	Q33782	dasyurus al
35	98	5.1	381	1	CYB_DASAL	Q34289	dasyurus al
36	98	5.1	381	1	CYB_SWICR	P55781	smnthopsis
37	98	5.1	460	1	NU4N_GADMO	P55781	gadus morhua
38	98	5.1	217	1	YPRA_ECOLI	P13974	escherichia
39	97.5	5.1	381	1	CYB_DASHA	O34311	dasyurus ba

Db 191 VAVDYITVALLINFGVGVGMIAHWRKPLRLQQAQYLMISALMALVFIKYLPEWTAWLIL 250
QY 181 GATSVYDLVAVLCCKPLRLMVLTAQERNEPFPALITYSSAMVTVGMKLDPSOGALQ 240
Db 251 AVISYDLVAVLCCKPLRLMVLTAQERNEPFPALITYSSAMVTVGMKLDPSOGALQ --RR 308
QY 241 LPYDP-----EMEEDSYSGFSPSPYEPFPLTGYPG-----EEL--- 277
Db 309 VPKNPYSTOGTERETQDTGTGSDGGSEWEAQDRLHGHPRSTPSRAVQELSGS 368
QY 277 -----EEDERGKVLGDFIFYSVLVKGAAATGSGDWNTTACFAVAILGLCTLALLAV 332
Db 369 ILTSEDEPERGVKVLGDFIFYSVLVKGASATASGDWNTTACFAVAILGLCTLALLAI 428
QY 333 FKALPALPISITFGILFYFSTDLNLRPMDTLASHQLYI 372
Db 429 FKALPALPISITFGILFYFATDYLQVPMQDLAFHQFYI 468
RESULT 9
PSNL_HUMAN
ID PSNL_HUMAN STANDARD; PRT: 467 AA.
AC P49768; Q14762; Q15719; Q15720;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN 1 (PS-1) (S182 PROTEIN).
GN PSNL OR PSNL1 OR AD3 OR PSI.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.; AND VARIANTS AD (FORMS I-467 AND I-463).
RC TISSUE=BRAIN;
RX MEDLINE; 95319502.
RA SHERRINGTON R., ROGAEV E.I., LIANG Y., ROGAEVA E.A., LEVESQUE G.,
RA IKEDA M., CHI H., LIN C., HOLMAN K., TSUDA T., MAR L.,
RA FONGIN J.-F., BRUNI A.C., MONTESI M.P., SORBI S., RAINERO I.,
RA PINESSI L., NEE L., CHUMAKOV I., POLLEN D., BROOKES A.,
RA SANSEAU P., POLINSKY R.J., WASSO W., DA SILVA H.A.R., HAINES J.L.,
RA PERICAK-VANCE M.A., TANZI R.E., ROSES A.D., FRASER P.E.,
RA ROMMENS J.M., ST GEORGE-HYSLOP P.H.;
RT "Cloning of a gene bearing missense mutations in early-onset familial
RT Alzheimer's disease.";
RL Nature 375:754-760(1995).
RN [2]
RP SEQUENCE FROM N.A. (FORMS I-463 AND I-374).
RC TISSUE=BLOOD, AND BRAIN;
RX MEDLINE; 96193901.
RA SAHARA N., YAHAGI Y.-I., TAKAGI H., KONDO T., OKOCHI M., USAMI M.,
RA SHIRASAWA T., MORI H.;
RT "Identification and characterization of presenilin I-463 and
RT I-374.";
RL FEBS Lett. 381:7-11(1996).
RN [3]
RP SEQUENCE OF 1-113 FROM N.A.
RA TSUJIMURA A., HASHIMOTO-GOTOH T.;
RL Submitted (MAR-1996) to the EMBL/GenBank/DBJ databases.
RN [4]
RP REVIEW ON VARIANTS.
RX MEDLINE; 98180715.
RA CRUTS M., VAN BROECKHOVEN C.;
RT "Presenilin mutations in Alzheimer's disease.";
RL Hum. Mutat. 11:183-190(1998).
RN [5]
RP VARIANTS AD THR-143 AND ALA-384.
RX MEDLINE; 96177673.
RA CRUTS M., BACKHOVENS H., WANG S.-Y., VAN GASSEN G., THEUNS J.,
RA DE JONGHE C., WEHRT A., DE VOECHT J., DE WINTER G., CRAS P.,
RA BRUYLAND M., DATSON N., WEISSENBAACH J., DEN DUNNEN J.T., MARTIN J.-J.,
RA HENDRIKS L., VAN BROECKHOVEN C.;
RT "Molecular genetic analysis of familial early-onset Alzheimer's
RT disease linked to chromosome 14q24.3.";

Hum. Mol. Genet. 4:2363-2372(1995).
RN [6]
RP VARIANTS AD L-82; H-115; T-139; R-163; T-231; L-264; V-392 AND Y-410.
RX MEDLINE; 96177674.
RA CAMPION D., FLAMAN J.-M., BRICE A., HANNEQUIN D., DUBOIS B.,
RA MARTIN C., MOREAU V., CHARBONNIER F., DIDIERJEAN O., TARDIEU S.,
RA PENET C., PUEL M., PASQUET F., LE DOZE F., BELLIS G., CALEND A.,
RA HEILIG R., MARTINEZ M., MALLET J., BELLIS M., CLERGET-DARPOUX F.,
RA AGID Y., FREBOURG T.;
RT "Mutations of the presenilin I gene in families with early-onset
RT Alzheimer's disease.";
RL Hum. Mol. Genet. 4:2373-2377(1995).
RN [7]
RP VARIANTS AD VAL-260; VAL-285 AND VAL-392.
RX MEDLINE; 95379971.
RA ROGAEV E.I., SHERRINGTON R., ROGAEVA E.A., LEVESQUE G., IKEDA M.,
RA LIANG Y., CHI H., LIN C., HOLMAN K., TSUDA T., MAR L., SORBI S.,
RA NACMIAS B., PIACENTINI S., AMADUCCI L., CHUMAKOV I., COHEN D.,
RA LANFELT L., FRASER P.E., ROMMENS J.M., ST GEORGE-HYSLOP P.H.;
RT "Familial Alzheimer's disease in kindreds with missense mutations in
RT a gene on chromosome 1 related to the Alzheimer's disease type 3
RT gene.";
RL Nature 376:775-778(1995).
RN [8]
RP VARIANTS AD V-139; V-146; Y-163; T-267; A-280 AND G-280.
RX MEDLINE; 96024664.
RA CLARK R.F., HUTTON M., FULDNER R.A., FROELICH S., KARRAN E.,
RA TALBOT C., CROOK R., LENDON C., PRIHAR G., HE C., KORENBLAT K.,
RA MARTINEZ A., WRAGG M., BUSFIELD F., BEHRENS M.I., MYERS A., NORTON J.,
RA MORRIS J., MEHTA N., PEARSON C., LINCOLN S., BAKER M., DUFF K.,
RA ZEHRE C., PREZ-TUR J., HOULDEN H., RUIZ A., OSSA J., LOPERA F.,
RA ARCOS M., MADRIGAL L., COLLINGE J., HUMPHREYS C., ASWORTH T.,
RA SARNER S., FOX N., HARVEY R., KENNEDY A., ROQUES P., CLINE R.T.,
RA PHILLIPS C.A., VENTER J.C., FORSEL L., AXELMAN K., LILIUS L.,
RA JOHNSTON J., COMBURN R., VIITANEN M., WINBLAD B., KOSIK K., HALTIA M.,
RA POYHONEN M., DICKSON D., MANN D., NEARY D., SNOWDEN J., LANTOS P.,
RA LANFELT L., ROSSOR M., ROBERTS G.W., ADAMS M.D., HARDY J., GOATE A.;
RT "The structure of the presenilin 1 (S182) gene and identification of
RT six novel mutations in early onset AD families. Alzheimer's Disease
RT Collaborative Group.";
RL Nat. Genet. 11:219-222(1995).
RN [9]
RP VARIANTS AD PHE-96; ARG-163 AND THR-213.
RX MEDLINE; 96310408.
RA KAMINO K., SATO S., SAKAKI Y., YOSHIIWA A., NISHIWAKI Y., TAKEDA H.,
RA TANABE H., NISHIMURA T., LI K., ST GEORGE-HYSLOP P.H., MIKI T.,
RA OGIHARA T.;
RT "Three different mutations of presenilin 1 gene in early-onset
RT Alzheimer's disease families.";
RL Neurosci. Lett. 208:195-198(1996).
RN [10]
RP VARIANT AD ALA-280.
RX MEDLINE; 97442268.
RA LENDON C.L., MARTINEZ A., BEHRENS I.M., KOSIK K.S., MADRIGAL L.,
RA NORTON J., NEUMAN R., MYERS A., BUSFIELD F., WRAGG M., ARCOS M.,
RA ARANGO VIANA J.C., OSSA J., RUIZ A., GOATE A.M., LOPERA F.;
RT "E280A PS-1 mutation causes Alzheimer's disease but age of onset is
RT not modified by ApoE alleles.";
RL Hum. Mutat. 10:186-195(1997).
RN [11]
RP VARIANTS AD THR-233 AND THR-278.
RX MEDLINE; 97316242.
RA KWOK J.B.J., TADDEI K., HALLUPP M., FISHER C., BROOKS W.S., BROE G.A.,
RA HARDY J., FULHAM M.J., NICHOLSON G.A., STELL R.,
RA ST GEORGE-HYSLOP P.H., FRASER P.E., KAKULAS B., CLARINETTE R.,
RA RELKIN N., GANDY S.E., SCHOFIELD P.R., MARTINS R.N.;
RT "Two novel (M23T and R278T) presenilin-1 mutations in early-onset
RT Alzheimer's disease pedigrees and preliminary evidence for
RL association of presenilin-1 mutations with a novel phenotype.";
RN NeuroReport 8:1537-1542(1997).
RP VARIANT GLY-318.
RX MEDLINE; 99115106.

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CC EMBL; Z71333; CA95930.1; -;
CC PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Alternative splicing; Glycoprotein.
FT TRANSMEM 83 103 POTENTIAL.
FT TRANSMEM 133 153 POTENTIAL.
FT TRANSMEM 161 181 POTENTIAL.
FT TRANSMEM 191 211 POTENTIAL.
FT TRANSMEM 221 241 POTENTIAL.
FT TRANSMEM 244 264 POTENTIAL.
FT TRANSMEM 281 301 POTENTIAL.
FT TRANSMEM 408 428 POTENTIAL.
FT TRANSMEM 433 453 POTENTIAL.
FT CARBOHYD 279 279 POTENTIAL.
FT CARBOHYD 405 405 POTENTIAL.
FT VARSPPLIC 26 29 MISSING (IN ISOFORM I-463).
SQ SEQUENCE 467 AA; 52384 MW; A841A0B7 CRC32;

Query Match 73.5%; Score 1413.5; DB 1; Length 467;
Best Local Similarity 71.0%; Pred. No. 7.2e-90;
Matches 282; Conservative 34; Mismatches 56; Indels 25; Gaps 4;

Qy 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRL 60
Db 71 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRL 130
Qy 61 NSVLTLMISVIVVMTIFLVLYKRYCYKFTGHWLIMSSLMMLFLFYIYVIGELVKYN 120
Db 131 HSNLNAALMISVIVVMTIFLVLYKRYCYKFTGHWLIMSSLMMLFLFYIYVIGELVKYN 190
Qy 121 VAMDYPTLLTWNFGAVGVCIIHWKGPLVLQOAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 191 VAVDYITVALLIMNFGVVGMSIHWKGPLRLQOAYLIMISALMALVFIKYLPEWTAWLIL 250
Qy 181 GAISYDVLAVLCPKGPLRMLVETAEQNERPIFPALIIYSSAMVTVGMAKLDPPSQGAL- 240
Db 251 AVISYDVLAVLCPKGPLRMLVETAEQNERPIFPALIIYSSAMVTVGMAKLDPPSQGAL- 310
Qy 240 -QLPYD-----PEMEEDSYDSFGSPSYPEVPEP-----PLTGYGPEEL----- 277
Db 311 KNTYNAOCTEREAQSPVNDGGFSEWEAQRDSQGLPHRSVTSVRAAQVSESSIPA 370
Qy 277 -EEEEERGVKGLGDFIFYSVLVGKAATGSDWNTTILACFVAILIGLCLTLLLAFFKK 335
Db 371 SEDPEERGKVLGDFVIFYSVLVGKASATASGDWNTTILACFVAILIGLCLTLLLAFFKK 430
Qy 336 ALPALPISITGLFIYFSTDNLRPFMDTLASHQYLI 372
Db 431 ALPALPISITGLFIYFATDYLVOPMDQLAFHQFYI 467

RESULT 11

PSN_DROME STANDARD; PRT; 541 AA.
AC 002194; 002395; 076802;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN HOMOLOG (DPS) (DMPS).
GN PS.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RP SEQUENCE FROM N.A. (LONG ISOFORM).
RX MEDLINE; 97285868.
RA BOULIANNE G.L., LYNE-BAR I., HUMPHREYS J.M., LIANG Y., LIN C.,
RA ROGAEV E., ST GEORGE-HYSLOP P.;
RT "Cloning and characterization of the Drosophila presenilin

homologue.";
RL NeuroReport 8:1025-1029(1997).
RN [2]
RP SEQUENCE FROM N.A. (SHORT ISOFORM).
RC STRAIN-CANTON-S;
RA HONG C.S., KOO E.H.;
RL "Isolation and characterization of Drosophila presenilin homolog.";
RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A. (LONG AND SHORT ISOFORMS).
RA YE Y., FORTINI M.E.;
RL "Characterization of Drosophila presenilin and its colocalization
with Notch during development.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC -----
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CC -----
DR EMBL; U77934; AAB61139.1; -;
DR EMBL; U78084; AAB53369.1; -;
DR EMBL; AF084184; AAC33129.1; -;
DR EMBL; AF084184; AAC33128.1; -;
DR FLYBASE; FBgn0019947; PS.
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Glycoprotein; Alternative splicing.
FT TRANSMEM 107 127 POTENTIAL.
FT TRANSMEM 155 175 POTENTIAL.
FT TRANSMEM 183 203 POTENTIAL.
FT TRANSMEM 217 237 POTENTIAL.
FT TRANSMEM 243 263 POTENTIAL.
FT TRANSMEM 266 286 POTENTIAL.
FT TRANSMEM 304 324 POTENTIAL.
FT TRANSMEM 482 502 POTENTIAL.
FT TRANSMEM 507 527 POTENTIAL.
FT CARBOHYD 129 129 POTENTIAL.
FT CARBOHYD 339 339 POTENTIAL.
FT CARBOHYD 410 410 POTENTIAL.
FT VARSPPLIC 384 397 MISSING (IN SHORT ISOFORM).
FT CONFLICT 80 81 GG -> RS (IN REF. 2).
SQ SEQUENCE 541 AA; 59304 MW; 796C4FE0 CRC32;

Query Match 58.6%; Score 1127.5; DB 1; Length 541;
Best Local Similarity 50.8%; Pred. No. 2.9e-70;
Matches 231; Conservative 61; Mismatches 74; Indels 89; Gaps 7;

Qy 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRL 60
Db 93 EEQGLKYGAQHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRL 152
Qy 61 NSVLTLMISVIVVMTIFLVLYKRYCYKFTGHWLIMSSLMMLFLFYIYVIGELVKYN 120
Db 153 SALANSLIMSVVMTIFLVLYKRYCYKFTGHWLIMSSLMMLFLFYIYVIGELVKYN 212
Qy 121 VAMDYPTLLTWNFGAVGVCIIHWKGPLVLQOAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 213 IPMDYPTALLIMNFGVVGMSIHWKGPLRLQOAYLIMISALMALVFIKYLPEWTAWL 272
Qy 181 GAISYDVLAVLCPKGPLRMLVETAEQNERPIFPALIIYSSAMVTV-----GMK 230
Db 273 AAIISWDLIAVLSPRGLRILVETAEQNERPIFPALIIYSSAMVTV-----GMK 332
Qy 231 LDPSS-----QGALQLP-----YDPEMEEDSYDSFGSPSYPEV-FEP 266
Db 333 SSPSSNSTTTTTRATONSLASPEAAASQRTGNSHPRONRDGGSVLATEGMLPTFKS 392

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Query Match          50.58; Score 972; DB 1; Length 461;
Best Local Similarity 52.88; Pred. No. 1e-59;
Matches 211; Conservative 53; Mismatches 74; Indels 62; Gaps 9;

QY 1 BELTKYGAHVIMLFVPTVLCMIVVATIKSVRFYEKNGQ-LIYPTFTEDTSPVGQRL 59
DB 39 EEAEKYGASHVHLFPVSLCMALVFTNTITFYSONNGRHLLYTPFVRETDSIVEKG 98

QY 60 LNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLLFTYIYLGVLKTY 119
DB 99 LMSGNALVMLCVVMTVLLVIFYKFKYKLIHGWLIVSSFLFLFTTIYVQEVLSKF 158

QY 120 NVAMDYPTLLTVNFGAVGMCVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVI 179
DB 159 DVSPSALLVLFGLCNYGVGLGMMCIHWKGPLRLQQLFYLLTMSALMALVFIKYLPEWTYVWF 218

QY 180 LGATSVYDLVAVLCPKGLPLMVLTAERNEPIPPALYISSANVYVGMKADPSSQOAL 239
DB 219 LFVSIWDLVAVLTPKGLRYLVTAEARNEPIPPALYISSGVYYPVLVT-----AV 271

QY 240 QLPYDPEWEDSYDF-----GP-----PSYPEVFEPP-----LTGYPE 274
DB 272 ENTTPR-EPITSSDNTSTAPPGEASCSSETPKPKWKRIPIKVOIESNTASTTQNSGV 330

QY 275 ELE-----EEERGKVLGLGDFIYFVSLVGRKAAATGSDWNTTACF 316
DB 331 RVERELAAERTVODANFHRHEERGKVLGLGDFIYFVSLVGRKAS--YEDWNTTIACY 388

QY 317 VAILIGLCITLLLVAPKKALPALPISITFGLIFYSTDN 356
DB 389 VAILIGLCITLLVAVFKRALPALQ-----PPFSPDS 420

RESULT 14
HOPI_CAEEL STANDARD; PRT; 358 AA.
ID HOPI_CAEEL AC 002100;
AC 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE INTEGRAL MEMBRANE PROTEIN HOP-1.
GN HOP-1 OR C18E3.8.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
SEQUENCE FROM N.A.
RC STRAIN-BRISTOL N2;
RC MEDLINE; 98004548.
RA LI X., GREENWALD I.;
RT "HOP-1, a Caenorhabditis elegans presenilin, appears to be
RT functionally redundant with SEL-12 presenilin and to facilitate LIN-12
RT and GLP-1 signaling.";
RL Proc. Natl. Acad. Sci. U.S.A. 94:12204-12209(1997).
RN [2]
SEQUENCE FROM N.A.
RC STRAIN-BRISTOL N2;
RC CONNELL M., MAGGI L.;
RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: MAY FACILITATE LIN-12 AND GLP-1 MEDIATED RECEPTION OF
CC INTERCELLULAR SIGNALS.
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
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OM protein - protein search, using sw model

Run on: March 18, 2000, 22:07:43 ; Search time 49.26 Seconds
(without alignments)
523.595 Million cell updates/sec

Title: US-08-509-359B-138
Perfect score: 1923
Sequence: 1 EELTLKYGAKHVIMLFVPT.....STDNLVRFPMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 225878 seqs, 69334122 residues

Database : SPTREMBL_12.*

Word size : 0

Number of hits that pass the threshold : 225878

- 1: sp.archaea.*
- 2: sp.bacteria.*
- 3: sp.fungi.*
- 4: sp.human.*
- 5: sp.invertebrate.*
- 6: sp.mammal.*
- 7: sp.mhc.*
- 8: sp.organelle.*
- 9: sp.phage.*
- 10: sp.plant.*
- 11: sp.rodent.*
- 12: sp.virus.*
- 13: sp.vertebrate.*
- 14: sp.unclassified.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	1896	98.6	449	6 Q9XT96	Q9xt96 bos taurus
2	1411.5	73.4	478	6 Q9XT97	Q9xt97 bos taurus
3	1398	72.7	456	13 Q9W6T7	Q9w6t7 brachydanio
4	1355	70.5	384	13 Q73869	O73869 cyprinus ca
5	576.5	30.0	272	5 O96340	O96340 drosophila
6	402.5	20.9	184	4 O95465	O95465 homo sapien
7	113.5	5.9	406	5 Q19737	Q19737 caenorhabdi
8	112	5.8	4578	13 Q42181	Q42181 fugu rubrip
9	110.5	5.7	320	8 Q34086	Q34086 coccyzus er
10	110.5	5.7	381	8 Q35425	Q35425 phascosor
11	108.5	5.6	380	8 Q922C9	Q922c9 upupa epops
12	107.5	5.6	748	2 Q92577	Q92577 streptomyc
13	107	5.6	381	8 Q3723	Q3723 antechinus
14	106.5	5.5	382	8 Q34340	Q34340 didelphis m
15	106.5	5.5	318	11 P97829	P97829 rattus norv
16	106.5	5.5	303	11 Q35294	Q35294 rattus norv
17	106	5.5	652	5 Q93346	Q93346 caenorhabdi
18	106	5.5	382	8 Q34279	Q34279 didelphis a
19	106	5.5	382	8 Q34677	Q34677 glirophia ve
20	105	5.5	444	2 Q9X2N3	Q9x2n3 arthrobacte
21	105	5.5	379	8 Q34428	Q34428 echinys did
22	104	5.4	382	8 Q35561	Q35561 philander o
23	104	5.4	379	8 Q36096	Q36096 trinomys pa
24	104	5.4	379	8 Q34430	Q34430 echinys did
25	104	5.4	826	10 O80739	O80739 arabidopsis

ALIGNMENTS

RESULT 1

Q9XT96 PRELIMINARY; PRT; 449 AA.
AC Q9XT96;
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-NOV-1999 (TREMBlrel. 12, Last sequence update)
DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
DE PRESENILIN 2.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovinae; Bos.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-BRAIN;
RA SAHARA N., SHIRASAWA T., MORI H.;
RT "Molecular cloning of bovine presentin 2 gene."
RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF038937; AAD39024.1; -
SQ SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;

Query Match 98.6%; Score 1896; DB 6; Length 449;
Best Local Similarity 98.4%; Pred. No. 9.9e-131;
Matches 366; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 EELTLKYGAKHVIMLFVPTLCMIVVATIKSVREYTEKNGOLIYTPFTEDPSVGORLL 60
|||||
Db 78 EELTLKYGAKHVIMLFVPTLCMIVVATIKSVREYTEKNGOLIYTPFTEDPSVGORLL 137
|||||
Qy 61 NSVLTNLMISVIVVMTIFLVVLYKYRCYKFTHGWLIMSSMLLFLFTYIYGEVLKTYN 120
|||||
Db 138 NSVLTNLMISVIVVMTIFLVVLYKYRCYKFTHGWLIMSSMLLFLFTYIYGEVLKTYN 197
|||||
Qy 121 VAMDYPTLLIWNFGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLEWSAWVL 180
|||||
Db 198 VAMDYPTLLIWNFGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLEWSAWVL 257
|||||
Qy 181 GAISYDVLAVLCPKGLRMLVETAQERNEFFPALIYSSAMVWTVGMKLPSSOGALQ 240
|||||
Db 258 GAISYDVLAVLCPKGLRMLVETAQERNEFFPALIYSSAMVWTVGMKLPSSOGALQ 317
|||||
Qy 241 LPYDPEMEEDSYDSFGSPSEYVFPPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 300
|||||
Db 318 LPYDPEMEEDSYDSFGSPSEYVFPPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 377
|||||
Qy 301 AAATGSGDNTLACFAVAILIGLCITLLLLAVFKKALPALPISITFGLIFYSTDNLVRP 360
|||||

Db 1 IMLFIPVTLKVVVVVATIKSVSYTKDQGOQLIYPPFREDTETVQGRALNSMLNAAIMIS 60
Qy 72 VIVVMTIFLVLYKYRCYKFIHGLWIMSLMLLFTYIYLGELVKTNVAMDYPTLLLT 131
Db 61 VIVVMTLVVLYKYRCYKFIHGLWIMSLMLLFTYIYLGELVKTNVAMDYPTLLLT 120
Qy 132 VVNFAGVGVCHHKRGPLVLAQYALIMISALMALVFIKYLPEWSAWILGALSVDLVAV 191
Db 121 IWNFGVGVCHHKRGPLVLAQYALIMISALMALVFIKYLPEWTAWILGALSVDLVAV 180
Qy 192 LCPKGPLRMLVETAQERNEPIFALIYSSAMVTVGMA-KLDPSQOGLQLP----- 243
Db 181 LCPKGPLRILVETAQERNEPIFALIYSSAMVTVGMA-KLDPSQOGLQLP----- 240
Qy 243 -----YDPMEDSDVSDFG-----EPSYDEVEPEPTGYPGBELEEEERGKVL 286
Db 241 APTAQPEDGGFTPAWVNOQHQGLQPMOSTEDSRREIQELPSARPP--PVEDDEERGKVL 298
Qy 287 GLGDFIFSVLVGKAAATGSGDWNNTLACFVAILGLCLTLLLLAVFKKALPALPISITF 346
Db 299 GLGDFIFSVMLVGRASATASGDWNNTIACFVAILGLCLTLLLLAIFKALPALPISITF 358
Qy 347 GLIFYSTDNLVRRPMDTLASHQLYI 372
Db 359 GLVFFVATDNLVRRPMDQLAVHQFYI 384

RESULT 5
O96340 ID O96340 PRELIMINARY; PRT; 272 AA.
AC O96340;
DT 01-MAY-1999 (TReMBLrel. 10, Created)
DT 01-MAY-1999 (TReMBLrel. 10, Last sequence update)
DE PRESENILIN (FRAGMENT).
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
RN Ephydroidea; Drosophilidae; Drosophila.
RP SEQUENCE FROM N.A.
RC STRAIN=CANTON S;
RX MEDLINE; 98331525.
RA MARFANY G., DEL-FAVERO J., VALERO R., DE JONGHE C., WOODROW S.,
RA HENDRIKS L., VAN BROECKHOVEN C., GONZALEZ-DUARTE R.;
RT "Identification of a Drosophila presenilin homologue: evidence of
RT alternatively spliced forms.";
RL J. Neurogenet. 12:41-54(1998).
DR EMBL; AF017025; AAD01611.1; -.
ET NON_TER
SQ SEQUENCE 272 AA; 29456 MW; 606B9A5C CRC32;

Query Match 30.0%; Score 576.5; DB 5; Length 272;
Best Local Similarity 47.8%; Pred. No. 5.8e-35;
Matches 133; Conservative 29; Mismatches 41; Indels 75; Gaps 8;
Qy 164 ALVFIKYLPWSAWILGALSVDLVAVLCPKGPLRMLVETAQERNEPIFALIYSSAMV 223
Db 1 ALVFIKYLPWTAWAVLAASISWDLIAVLSPRGLRILVETAQERNEQIFALIYSSIVV 60
Qy 224 WTV-----GMAKLDPS-----QGALQLPYDPEMEEDSDVSDFGEPSPYE- 263
Db 61 YALVNTVTPQSOATASSPSSSSNTTTTRATQNSLA---SPEAAAASQRTGN-SHPRQ 116
Qy 263 -----VREPLTGYPGE---ELEE----- 279
Db 117 NORDGSLVATEAEAGTFQWSANLSERVARRQLEVOSTQSGNAQRSENEYFTVTPDON 176
Qy 279 ----EEERGKVLGLGDFIFSVLVGKAAATGSGDWNNTLACFVAILGLCLTLLLLAVFK 334
Db 177 HPDGOEERGKVLGLGDFIFSVLVGKASS--YGDWTTIACFVAILGLCLTLLLLATWR 234

Qy 335 KALPALPISITGLIFYSTDNLVRRPMDTLASHQLYI 372
Db 235 KALPALPISITGLIFCFEATSAVVPFPMEDLSAKQVFI 272
RESULT 6
O95465 ID O95465 PRELIMINARY; PRT; 184 AA.
AC O95465;
DT 01-MAY-1999 (TReMBLrel. 10, Created)
DT 01-MAY-1999 (TReMBLrel. 10, Last sequence update)
DE MINILIN.
GN PSNI.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RA POWELL C.S., GEGG M.E., PALMER M.S.;
RT "Human presenilin 1 gene encodes an alternative protein-minilin.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DDBJ databases.
DR EMBL; AJ008005; CAA07825.1; -.
SQ SEQUENCE 184 AA; 21073 MW; 5C6FBAEE CRC32;

Query Match 20.9%; Score 402.5; DB 4; Length 184;
Best Local Similarity 73.5%; Pred. No. 2.1e-22;
Matches 83; Conservative 9; Mismatches 16; Indels 5; Gaps 1;
Qy 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFTEKNGQLIYPTFTDTPSVGORLL 60
Db 71 EELTKYGAHVIMLFVPTLCMIVVATIKSVFSYTRKDGQLIYPTFTDTPSVGORAL 130
Qy 61 NSVLNTLMISVIVVMTIFLVLYKYRCYKFIHGLWIMSLMLLFTYIYLG 113
Db 131 HSILNAINISVIVVMTILLVLYKYRCYK----VSMRHSLLSLFLFWLG 178

RESULT 7
Q19737 ID Q19737 PRELIMINARY; PRT; 406 AA.
AC Q19737; Q22692;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-MAY-1999 (TReMBLrel. 10, Last sequence update)
DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
DE F22E10.5 PROTEIN.
GN F22E10.5.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
OC Rhabditina; Rhabditoidae; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
RP SEQUENCE FROM N.A.
RA GARDNER A.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DDBJ databases.
DR EMBL; Z67882; CAA91804.1; -.
DR EMBL; Z50797; CAA91804.1; JOINED.
DR EMBL; Z50797; CAA90677.1; -.
DR EMBL; Z67882; CAA90677.1; JOINED.
DR PROSITE; PS00379; CDP_ALCOHOL_P_TRANSF; 1.
SQ SEQUENCE 406 AA; 45628 MW; 07336492 CRC32;

Query Match 5.9%; Score 113.5; DB 5; Length 406;
Best Local Similarity 24.7%; Pred. No. 0.54;
Matches 64; Conservative 31; Mismatches 71; Indels 93; Gaps 13;
Qy 7 YGAKHVIMLFVPTLCM-----IYVATIKSVRFY-----TEKNGQLIYPTF--T 49
Db 136 HGCDSMTQVFTLNICYAMSLGTPVGVGLIVSVIVMVFCAHWSYCTGQLRFSKFDVT 195
Qy 50 EDTPSPVGORLL-----NSVLNTLI-----MISVIVVMTIFLVLYKYRCYKFIHGLW 97

Search completed: March 18, 2000, 22:07:45
Job time: 125 sec

Query Match 5.5%; Score 106.5; DB 8; Length 382;
Best Local Similarity 20.5%; Pred. No. 1.6;
Matches 79; Conservative 53; Mismatches 115; Indels 139; Gaps 18;

QY 21 LCMIVVATIKSVRFYTEKNGQLIYPTFTEDTPSVGQRLNSVLNTLIMISVIVVMTIFL 80
Db 39 MCLLIQILT-----GLFLAMHYTSDT-----LTAFS 64
QY 81 VVLYKYRCYFIHGLIM-----SSLMLLFYIYIYGLVLYKTYNVAMDYPTLLLTWN 134
Db 65 SV--AHICRDVNYGLIRNIHANGASFFMCLFLHVGRIYIGSY-----LYKETWN 114
QY 135 FGAVGVCIHWKGPLVLQQAAYLIMISALMALVFIKYIPEWSAWILGAIISVYDLVAVLCP 194
Db 115 IG-----VILLVFMATAFVGYYLPMGQMSFGWATVITNLLSAIPY 155
QY 195 KGPLRLMVE-----TAQERNEPIFPALYISSAMVWTVMGAK-----LDP 233
Db 156 IG--NTLVEWINGGFSVDKATLIRFFAFHFILFPIILAMVVHLLFHERGSNNPTGLDP 213
QY 234 SSQCALQLPYDP-----EMEDSYDSFGSPSYPEVFEP--PLTGYPG 273
Db 214 NSD---KIPHPYTYTIKDILGLFLMIILLSLAMFSPDLLGD---PDNFTPANLNTPPH 267
QY 274 EELEEEERGVKLGDFIF-YSVLVGKAAATGSGDNTTILACFVAILIGLCLTLLLLAV 332
Db 268 IKPE-----WYFLPAYAILRIPNKLGG-----VLALLASILILLPLLHTST 311
QY 333 FKALPALPISITFGLIFYESTDNLV 358
Db 312 -QRSMFRPISQT---LFWMLTANLI 333

RESULT 15

O35294
ID O35294 PRELIMINARY; PRT; 318 AA.
AC O35294;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-MAY-1999 (TREMBLrel. 10, Last annotation update)
DE INTEGRIN-ASSOCIATED PROTEIN FORM 4.
GN IAP.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-SPRAGUE-DAWLEY; TISSUE-BRAIN;
RA HUANG A.M., LEE E.H.Y.;
RL Submitted (AUG-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF017437; AAB70273.1; -
KW Integrin.
SQ SEQUENCE 318 AA; 34776 MW; FAA078CB CRC32;

Query Match 5.5%; Score 106.5; DB 11; Length 318;
Best Local Similarity 23.0%; Pred. No. 1.3;
Matches 40; Conservative 35; Mismatches 50; Indels 49; Gaps 8;

QY 3 LTLKYGAKH-----VIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTFTEDTPSVGQ 57
Db 158 LTLKYKSSHTNKRILLVAGLALTIVV-----GAILFIP-----GE 196
QY 58 RLNSVLTNLTIMISVIVVMTIFLVLYKYCYKFIHGLWLMSSMLLFYIYVGLVK 117
Db 197 KPVKNASG-----LGLIVISTGIILL-QYNVFMTAFG---MTSFTIAILITQV-LGYVLA 247
QY 118 TYNVAMDYPTLLLTWNFGAVGVCIHWKGPLVLQQAAYLIMISALMALVFIKYL 171
Db 248 VVGWCLCI-----NACEPVHGPLLSGLIGLIALAELGLVYMKFV 287

OM of: US-08-509-359B-137 to: GenEmbl:* out_format : pfs

Date: Mar 18, 2000 2:37 PM

About: Results were produced by the GenCore software, version 4.5,
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Search information block:

Query: US-08-509-359B-137

Query length: 448

Database: GenEmbl.*

Database sequences: 821193

Database length: 1518192014

Search time (sec): 533.740000

score_list:

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gb_pr3:HUMES1P	+ 2336.00	3900.98	2.9e-209	2229	L44577 Homo sapiens S182 prote
gb_pr2:HUMSTM2R	+ 2328.00	3887.50	1.6e-208	2236	L43964 Homo sapiens (clone F-1
gb_om:AF038937	+ 2262.50	3778.58	1.9e-202	2004	AF038937 Bos taurus presenilin
gb_ro:MMU57324	+ 2232.00	3727.60	1.3e-199	1954	U57324 Mus musculus PS-2 mRNA,
gb_ro:AF038935	+ 2228.50	3724.54	2.0e-199	1450	AF038935 Mus musculus presenil
gb_ro:D83700	+ 2227.00	3723.07	2.4e-199	1347	D83700 Rattus norvegicus cDNA
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gb_ro:AB004454	+ 2213.00	3699.04	5.1e-198	1414	AB004454 Rattus norvegicus mRN
gb_ov:D84428	+ 1624.50	2704.30	1.3e-142	2450	D84428 Xenopus laevis mRNA for
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gb_ro:MMU57325	+ 774.00	1283.30	1.9e-63	1146	U57325 Mus musculus PS-2short

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gb_pl1:AP000836 - 567.00 882.27 4.0e-41 190014 ! AP000836 Oryza sativa gen
gb_htg4:AC009840 + 545.00 851.00 2.2e-39 109951 ! AC009840 Drosophila melan

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seq documentation block: 2285 bp DNA PAT 29-SEP-1999
LOCUS AR060156

DEFINITION Sequence 136 from patent US 5840540.

ACCESSION AR060156

VERSION AR060156.1 GI:5986606

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 2285)

AUTHORS St. George-Hyslop,P.H., Rommens,J.M. and Fraser,P.E.

TITLE Nucleic acids encoding presenilin II

JOURNAL Patent: US 5840540-A 136 24-NOV-1998;

FEATURES Location/Qualifiers

source

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BASE COUNT 537 a 579 c 633 g 521 t 15 others

ORIGIN

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Ratio: 5.214 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

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17 gThrSerLeuMetSerAlaGluSerProThrProArgSerCysGlnGlu 34

416 GACGTCCCTATATGCGCGGAGAGCCCGCCGCTCTGCGCAGGAGG 465

34 lyArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50

466 GCAGCAGGCGCCAGAGGATGAGAGATACTGCCAGCTGGAGAGCCAG 515

51 GluAsnGluGluAspGlyGluGluAspProAspArgTyrValCysSerG1 67

516 GAGAACGAGGAGGAGCGGTGAGGAGGACCTGACCGCTATGCTGTGTAGTG 565

67 yValProGlyArgProProGlyLeuGluGluGluLeuThrLeuLysTyrG 84

566 GTTCCCGGGCGCGCCGAGGAGGAGAGCTGACCTCAATAGG 615

84 lyAlaLysHisValIleMetLeuPheValProValThrLeuCysMetIle 100

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101 ValValValAlaThrIleLysSerValArgPheTyrThrGluLysAsnG1 117

666 GTGTGTGTAGCCACCATCAAGTCTGTGGCTTCTACACAGAGAGAAATGG 715

117 yGluLeuIleTyrThrProPheThrGluAspThrProSerValGlyGlnA 134

716 ACAGCTCATCTACAGCCCATTCATCTGAGGACACACCTCGGTGGCCAGC 765

134 rgLeuLeuAsnSerValLeuAsnThrLeuIleMetIleSerValIleVal 150

766 GCTTCCTCACTCGGTGTGACACCCCTCATCATGATCAGCGTCATCGTG 815

151 ValMetThrIlePheLeuValValLeuTyrLysTyrArgCysTyrLysph 167

JOURNAL Submitted (16-DEC-1997) Molecular Biology, Tokyo Institute of
Psychiatry, 2-1-8 Kami-Kitazawa, Setagaya-ku, Tokyo 156, Japan
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17 gThrSerLeuMetSerAlaGluSerProThrProArgSerCysGlnGlu 34
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133 GAGTCCCTGATGTACAGCGAGAGCCCTACGCCGCGCTCTGCCAGGACG 182
34 lYArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50
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183 GCCGGCAGGCGCTGGAGAGCGGAGAGTGCCTGCCAGTGGAGAGCCAG 232
51 GluAsnGluGluasp...GlyGluGluAspProAspArgTyrValCysSe 66
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233 GAGAGTGAGGAGGACACACGAGGAGGAGGACCCCTGACCGCTACGCTGCAG 282
66 rGlyValProGlyArgProGlyLeuGluGluGluLeuThrLeuLysT 83
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283 TGGGGTTCCTGGGCGCCCGAGGCTTGAGGAGGAGCTGACCTCAAT 332
83 yrcGlyAlaLysHisValIleMetLeuPheValProValThrLeuCysMet 99
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333 ACGGGCAAGCATGTGATGTTGTTGTGCTGTACACACTGTGCATG 382
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150 ValValMetThrIlePheLeuValValLeuTyrLysTyrArgCysTyrLy 166

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533 GTCACCATGACGATCTTCCTGGTGTGCTCTACAAGTACCGCTGCTACAA 582
166 sPheIleHisGlyTrpLeuIleMetSerSerLeuMetLeuLeuPheLeuP 183
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633 TCACCTACATCTACCTCGGGGAAGTGTCAAGACCTACAATGTGGCCATG 682
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683 GACTACCCACCCCTGTCTCGACCGTGTGAACCTCGGGCGGTGGGCAT 732
216 tValCysIleHisTrpLysGlyProLeuValLeuGlnGlnAlaTyrLeuI 233
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350 GluGluLeuGluGluGluGluGluGluGluGluGluGluGluGluGlu 366
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AUTHORS Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
JOURNAL 1 (bases 1 to 2088)
AUTHORS Frenzel, S., Abdel, A.S. and Luebbert, H.
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 2088)
AUTHORS Frenzel, S.
TITLE Direct Submission
JOURNAL Submitted (05-JUL-1996) S. Frenzel, Sandoz Pharma Ltd, Preclinical
RESEARCH, PO Box, CH-4002 Basel, Switzerland
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Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE 1 (bases 1 to 1673)
AUTHORS Li, J., Ma, J. and Potter, H.
TITLE Identification and expression analysis of a potential familial
Alzheimer disease gene on chromosome 1 related to AD3
Proc. Natl. Acad. Sci. U.S.A. 92 (26), 12180-12184 (1995)
JOURNAL 96109229
MEDLINE
REFERENCE 2 (bases 1 to 1673)
AUTHORS Li, J.
TITLE Direct Submission
JOURNAL Submitted (21-AUG-1995) Jinhe Li, Neurobiology, Harvard Medical
School, 220 Longwood Ave., B2-502, Boston, MA 02115, USA

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CDS

AUTHORS TITLE JOURNAL

Tanahashi, H.
Direct Submission
Submitted (28-MAY-1997) to the DDBJ/EMBL/GenBank databases. Hiroshi
Tanahashi, National Institute of Neuroscience, Division of
Demyelinating Disease and Aging; 4-1-1 Ogawahigashi, Kodaira, Tokyo
187, Japan (E-mail: tanahash@ncnp.nicp.go.jp, Tel: 81-423-41-1717,
Fax: 81-423-46-1747)

REFERENCE

2 (sites)

AUTHORS

Tanahashi, H. and Tabira, T.

TITLE

Cloning of the cDNA encoding rat presenilin-2
Biochim. Biophys. Acta 1396 (3), 259-262 (1998)

JOURNAL

98207716

MEDLINE

Sequence updated (06-Jun-1997)

COMMENT

Sequence updated (08-Jun-1997).

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911	TCGACTCAGCAGCGGTATCTCATATGATCAGTGCCCTCATGGCCCTGG	960
242	alPheIleLysTyrLeuProGluTrpSerAlaTrpValIleLeuGlyAla	258
961	TATTTATCAAGTACTCTCCCGAAATGGACCGCATGCTCATCTTGGCTGTG	1010
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1011	ATTTCAAGTATGATTTGGTGGCTGTTTTATGTCCCAAGGCCCACTTCG	1060
275	gMetLeuValGluThrAlaGlnGluArgAsnGluProIlePheProAlaI	292
1061	TATGCTGGTTGAACAGCTCAGGAAGAAATGAGACTCTCTTCTCAGCTC	1110
292	euleTyrSerSerAlaMetValTrpThrValGlyMetAlaLysLeuAsp	308
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159 LeuTyrLysTyrArqCysTyrLysPheIleHisGlyTrpLeuIleMetse 175
711 CTGTATAAAATACAGGTGCTACAAGGTTCATCCACGCGCTGCCTATTATTATTC 760
175 rSerLeuMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluVal 192
761 ATCTCTGTGTGTCTTCTTTTTCGTCATTACTTTAGGGAAGTAT 810
192 euIysThrTyrAsnValAlaMetAspTyrProThrLeuLeuLeuThrVal 208
811 TTAAGACCTACAATGTCCGCTGAGTACGTTACAGTACGACTCCCTAATC 860
209 TrpAsnPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProLe 235
861 TGGAAATTTGGTGTGGGTGGGATGANTGCCATCCACTGGAAAGGCCCT 910
225 uValLeuGlnGlnAlaTyrIleIleMetIleSerAlaLeuMetAlaLeuV 242
911 TCGACTGCAGCAGCGTATCTCATTATGATCAGTGCCTCATGGCCCTGG 960
242 alPheIleLysTyrLeuProGluTrpSerAlaTrpValIleLeuGlyAla 258
961 TATTTATCAAGTACTCTCCCGAATGGACCGCATGCTCATCTGGCTGTG 1010
259 IleSerValTyrAspLeuValAlaValLeuCysProLysGlyProLeuAr 275
1011 ATTTCAGTATGATTTTGGTGGCTGTTTTATGTCCCAAGGCCACTTCG 1060
275 gMetLeuValGluThrAlaGlnGluArgAsnGluProIlePheProAla 292
1061 TATGCTGGTTGAACAGCTCAGGAAGAAGTACAGACTCTCTTCTCAGCTC 1110
292 euIleTyrSerSerAlaMetValTrpThrValGlyMetAlaLysLeuAsp 308
1111 TTATCTATTTCTCAACATGTGTGGTTGGTGAATGTGCTCAAGGAGAC 1160

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292 eulleTyrSerSerAlaMetValTrpThrValGlyMetAlaIysLeuAsp 308
||||| | | | | | | |
|||||:::|||||||
|||||:::|||||||
1111 TTATCTATTCTCAACAATGGTGGTGGTGGTGAATGGCTGAAGGAGAC 1160

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COMMENT
FEATURES

OM of: US-08-509-359B-137 to: N_Geneseq_36:* out_format : pfs
Date: Mar 18, 2000 11:46 PM
About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-1998 CompuGen Ltd.

Command line parameters:
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-Q=/cgn2.1/USPTO_spool/US08509359/runat_17032000.164056.19497/app_query.fasta.1
-DB=N_Geneseq_36 -QFWT=fastap -SUFFIX=ring -GAPOP=12.000
-GAPEXT=4.000 -MINMATCH=0.100 -LOOPEXT=0.000 -LOOPEXT=0.000
-GAPOP=4.500 -GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500
-DELOP=6.000 -GAPEXT=7.000 -START=1 -MATRIX=blomsum62
-TRANS=human40.cdi -LIST=45 -DOCALLIGN=200 -THR_SCORE=pct
-ALIGN=15 -MODE=LOCAL -OUTFWT=pfs -NORM=ext -MINDEN=0
-MAXLEN=1000000 -USER=US08509359 -NCPU=6 -ICPU=3 -NO_XLPXY -WAIT
-THREADS=1

Search information block:
Query: US-08-509-359B-137
Query length: 448
Database: N_Geneseq_36:*
Database sequences: 311585
Database length: 125096042
Search time (sec): 37.450000

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N_Geneseq_36:T51253	+ 2336.00	4306.67	2.8e-232	2236	Human Ab4 protein coding sequ
N_Geneseq_36:V04669	+ 2336.00	4306.70	2.8e-232	2229	Human presenilin-2 cDNA (hps2)
N_Geneseq_36:T87426	+ 2320.50	4277.65	1.1e-230	2276	Full AD4/AD3LP sequence. Ident
N_Geneseq_36:T87401	+ 1623.50	2987.65	8.2e-159	1417	AD4/AD3LP sequence. Identifin
N_Geneseq_36:T40030	+ 1468.00	2694.82	1.7e-142	1964	Murine presenilin-1 wild type
N_Geneseq_36:T64819	+ 1468.00	2691.19	2.7e-142	2681	Tumour suppressor inhibited pa
N_Geneseq_36:V04668	+ 1468.00	2694.82	1.7e-142	1964	Mouse suppressor inhibited pa
N_Geneseq_36:T40028	+ 1467.00	2688.97	3.5e-142	2765	Presenilin-1-1 wild type codin
N_Geneseq_36:V17358	+ 1467.00	2688.98	3.5e-142	2764	PS1/467 protein coding sequen
N_Geneseq_36:V04666	+ 1467.00	2688.97	3.5e-142	2765	Human presenilin-1 cDNA (hps1-
N_Geneseq_36:T85332	+ 1462.00	2679.68	1.2e-141	2765	Human S182 gene, PS1 locus, re
N_Geneseq_36:V23525	+ 1460.00	2675.97	1.9e-141	2764	Homo sapiens PS-1 cDNA. Nucle
N_Geneseq_36:T59535	+ 1454.00	2670.06	4.0e-141	1762	Human early onset Alzheimer's
N_Geneseq_36:T59536	+ 1454.00	2670.14	4.0e-141	1750	Human early onset Alzheimer's
N_Geneseq_36:T75576	+ 1449.00	2659.80	1.5e-140	1914	Presenilin-1 VRQO variant gene
N_Geneseq_36:T40029	+ 1448.00	2652.38	3.9e-140	3086	Presenilin-1-2, alternatively
N_Geneseq_36:V04667	+ 1448.00	2652.38	3.9e-140	3086	Human presenilin-1 cDNA (hps1
N_Geneseq_36:V03246	+ 1442.00	2655.27	2.7e-140	1750	Human presenilin-1 cDNA. DNA e
N_Geneseq_36:T63207	+ 1442.00	2646.81	7.9e-140	1911	Human S182 gene associated wit
N_Geneseq_36:T87402	+ 1438.00	2642.29	1.4e-139	1488	Partial AD3 sequence. Identifi
N_Geneseq_36:V17357	+ 1438.00	2640.72	1.7e-139	1703	PS1/429 protein coding sequen
N_Geneseq_36:T85333	+ 1435.00	2629.50	7.3e-139	2765	Human mutant S182 gene, PS1 lo
N_Geneseq_36:T40043	+ 1166.50	2134.83	2.6e-111	1895	Presenilin homologue. New pres
N_Geneseq_36:T60306	+ 1003.50	1834.59	1.4e-94	1500	Caenorhabditis elegans SEL-12
N_Geneseq_36:T51271	+ 666.00	1220.72	2.1e-60	473	Human expressed sequence tag ES
N_Geneseq_36:T51258	+ 385.00	679.56	3.0e-30	2387	Human AD4 gene genomic sequen
N_Geneseq_36:T51256	+ 377.00	660.75	6.1e-28	2349	Human AD4 gene genomic sequen
N_Geneseq_36:X11281	+ 348.00	638.06	6.1e-28	230	Human biallelic polymorphic DNA
N_Geneseq_36:T40037	+ 315.00	560.25	1.3e-23	945	Human presenilin-1 gene exon 8
N_Geneseq_36:T99666	+ 315.00	560.25	1.3e-23	945	Human presenilin-1 gene exon 8
N_Geneseq_36:T51260	+ 270.00	459.79	5.2e-18	4004	Human AD4 gene genomic sequen
N_Geneseq_36:T51259	+ 262.00	452.67	1.3e-17	2058	Human AD4 gene genomic sequen
N_Geneseq_36:T51257	+ 229.50	396.44	1.8e-14	1438	Human AD4 gene genomic sequen
N_Geneseq_36:T40042	+ 224.00	394.02	2.4e-14	736	Presenilin-1 exon 13. New pres
N_Geneseq_36:T99671	+ 224.00	394.02	2.4e-14	736	Human presenilin-1 gene exon 13
N_Geneseq_36:T40034	+ 212.50	362.71	1.3e-12	1727	Presenilin-1 exon 5. New pres
N_Geneseq_36:T99663	+ 212.50	362.71	1.3e-12	1727	Human presenilin-1 gene exon 5
N_Geneseq_36:T40035	+ 201.00	340.33	2.3e-11	1893	Presenilin-1 exon 6. New pres
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N_Geneseq_36:T99670 + 191.00 329.08 9.9e-11 1003 ! Human presenilin-1 gene exo
N_Geneseq_36:T59473 + 186.00 319.87 3.2e-10 996 ! Early onset Alzheimer's dise
N_Geneseq_36:T40038 + 167.50 292.61 1.1e-08 540 ! Presenilin-1 exon 9. New pre
seq_name: N_Geneseq_36:T40031
seq_documentation_block:
ID T40031 standard; DNA; 2229 BP.
AC T40031;
DT 25-JUL-1997 (first entry)
DE Human presenilin-2 wild type coding sequence.
KW Presenilin-2; human; hps1-1; hps1-2; PS-2; integral membrane protein; AD;
KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;
KW depression; antibody; gene expression modulator; therapy; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
FT cds 366..1712
FT cds /*tag= a
FT FT /product= presenilin-2
FT FT
PN W09634099-A2.
PD 31-OCT-1996.
PF 29-APR-1996; CA0263.
PR 28-APR-1995; US-431048.
PR 28-JUN-1995; US-496841.
PR 31-JUL-1995; US-509359.
PA (HSCR-) HSC RES & DEV LP.
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
PI WPI; 96-497631/49.
PI P-PSDB: W05762.
PT New presenilin genes - useful for diagnosis, therapy and drug
PT screening of familial Alzheimer's disease, cerebral disorders, etc.
PS Claim 8; Page 148-150; 178pp; English.
CC This sequence represents the coding sequence for the human presenilin-2.
CC T40028 and T40029 represent the coding sequences for the two different
CC forms of wild type human presenilin-1 (PS-1). The form represented by
CC T40029 results from alternate splicing of the genomic DNA sequence.
CC T40030 represents the coding sequence for wild type mouse PS-1. The
CC presenilins are a family of highly conserved integral membrane proteins
CC with a common structural motif, common alternate splicing patterns, and
CC common mutational hot spot regions. Mutations in PS genes are implicated
CC in familial Alzheimer's disease (AD) and possibly other diseases such as
CC cerebral haemorrhage, schizophrenia, depression etc., so detection of
CC mutations in these sequences can be used for diagnosis of these diseases.
CC The encoded proteins, or vectors that express them or containing
CC antisense sequences, antibodies selective for mutant forms of the encoded
CC proteins (such as W05736) and modulators of PS gene expression are
CC potentially useful for treatment of AD etc. Transgenic animals are useful
CC as models for drug screening. The antibodies can also be used e.g. for
CC affinity purification and in immunoassays.
SQ Sequence 2229 BP: 481 A; 579 C; 633 G; 521 T;

alignment_scores:
Quality: 2336.00 Length: 448
Ratio: 5.214 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000
alignment_block:
US-08-509-359B-137 x T40031
Align seg 1/1 to: T40031 from: 1 to: 2229
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366 ATGCTCATTCATTCAGCTCTGTACAGCAGGAGGAAGTGTGTGATGAGCG 415
17 gThrSerLeuMetSerAlaGluSerProThrProArgSerCysGlnGluG 34
|||||
416 GACGTCCCTATATGTCGGCGAGAGCCACGCCGCGCTCTCCAGGAGG 465
34 lYArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50

CC acid probes specific for the mutant gene, provides a means of
 CC diagnosing Alzheimer's disease.

50 Sequence 2236 BP; 488 A; 584 C; 645 G; 519 T;

alignment_scores:

Quality: 2336.00 Length: 448
 Ratio: 5.214 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-08-509-359b-137 x T51253

Align seg 1/1 to: T51253 from: 1 to: 2236

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368 ATGCTCACATTCATGCGCCCTCAGACGAGGAGAAAGTGTGTATGAGCG 417
17 gThrSerLeuMetSerAlaGluSerProThrProArgSerCysGlnGlu 34
418 GAGTCCCTAAATGTGGCGGAGAGCCCGCCGCTCTGCCAGGAGG 467
34 lyArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50
468 GCAGGCAGGCGCCAGAGGATGGAGAAACACATGCCAGTGGAGAGCCAG 517
51 GluAsnGluGluAspGlyGluAspProAspArgTyrValCysSerGln 67
518 GAGAACGAGGAGGAGCGGTGAGGAGGACCTGACCGCTATGCTGTAGTGG 567
67 yValProGlyArgProProGlyLeuGluGluLeuThrLeuLysTyrG 84
568 GGTTCCTCCGGCGCGCCAGGCGTGGAGGAGAGCTGACCCCTCAAAATAG 617
84 lyAlaLysHisValIleMetLeuPheValProValThrLeuCysMetIle 100
618 GAGCGAAGCACGTGATCATGCTGTTTGTGCGTGTCACTGTGTGCATGATC 667
101 ValValValAlaThrIleLysSerValArgPheTyrThrGluLysAsnGln 117
668 GTGGTGTAGCCACCACCAAGCTGTGCGCTTCTACACAGAGAAGAATGG 717
117 yGlnLeuIleTyrThrProPheThrGluAspThrProSerValGlyGluA 134
718 ACAGCTCATCTACAGCCCAATCACTGAGGACACACCCCTCGGTGGGCGAG 767
134 rgLeuLeuAsnSerValLeuAsnThrLeuIleMetIleSerValIleVal 150
768 GCCTCTCACTCCGTCGTCGACACCCCTCATCATGATCAGCGTCATCGTG 817
151 ValMetThrIlePheLeuValValLeuTyrLysTyrArgCysTyrLysPh 167
818 GTTATGACCATCTTCTGTGTGTGTCCTACAAGTACCGCTGCTACAAGTT 867
167 erIleHisGlyTrpLeuIleMetSerSerLeuMetLeuLeuPheLeuPheT 184
868 CATCCATGCGCTGGTGTATCATGCTCTTCACTGATGCTGCTCTCTCTTCA 917
184 hrTyrIleTyrLeuGlyGluValLeuLysThrTyrAsnValAlaMetAsp 200
918 CCTATATCTACCTTGGGGAAGTGCTCAAGACCTTACAAATGTGGCCATGG 967
201 TyrProThrLeuLeuLeuThrValTrpAsnPheGlyAlaValGlyMetVa 217
968 TACCCACCCCTCTTCTGCTGACTGTCTGGAATTCGGGCGAGTGGCGATGG 1017
217 lCysIleHisTrpLysGlyProLeuValLeuGlnGlnAlaTyrLeuIleM 234
1018 GTGCATCCACTGGAAGGCGCCCTGTGTGTGTGTCAGCAGGCGCTACCTCA 1067
234 erIleSerAlaLeuMetAlaLeuValPheIleLysTyrLeuProGluTrp 250

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1068 TGATCAGTGGCGCTCATGGCCCTAGTGTTCATCAAGTACCTCCACAGATGG 1117
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267 lLeuCysProLysGlyProLeuArgMetLeuValGluThrAlaGlnGluA 284
1168 GCTGTGTCCTCCAAAGGGCCTCTGAGAATGCTGGTAGAAACTGCCCAGGAGA 1217
284 rgAsnGluProIlePheProAlaLeuIleTyrSerSerAlaMetValTrp 300
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301 ThrValGlyMetAlaLysLeuAspProSerSerGlnGlyAlaLeuGlnLe 317
1268 ACGGTTGGCATGGCAAGCTGGACCCCTCTCTCAGGGTGGCCCTCCAGCT 1317
317 uProTyrAspProGluMetGluGluAspSerTyrAspSerPheGlyGluP 334
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334 roSerTyrProGluValPheGluProProLeuThrGlyTyrProGlyGlu 350
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367 erIlePheTyrSerValLeuValGlyLysAlaAlaAlaThrGlySerGlyA 384
1468 CATCTTCTACAGTGTGCTGGTGGCAAGGCGGTGCCACGGGCGAGCGGG 1517
384 spTrpAsnThrThrLeuAlaCysPheValAlaIleLeuIleGlyLeuCys 400
1518 ACTGGAATACACGCTGGCCTGCTCTGCTGGCCATCCTCATTTGGTGTGT 1567
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1568 CTGACCCCTCTGCTGCTGTGTGTGTTCAAGAAGGCGCTGCCCGCCTCCC 1617
417 oIleSerIleThrPheGlyLeuIlePheTyrPheSerThrAspAsnLeuV 434
1618 CATCTCCATCACGTTGCGGCTCATCTTTTACTTCTCCACGGACACCTGG 1667
434 alArgProPheMetAspThrLeuAlaSerHisGlnLeuTyrIle 448
1668 TGGCGCGCTTCATGGACACCCCTGGCCCTCCCATCAGCTCTACATC 1711
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seq_documentation_block:
ID V04669 standard; CDNA; 2229 BP.
AC V04669;
DT 20-JUL-1998 (first entry)
DE Human presenilin-2 cDNA (hps2).
KW Presenilin-1; PS1 gene; human; familial Alzheimer's disease; FAD;
KW cerebral haemorrhage; schizophrenia; depression; epilepsy;
KW mental retardation; diagnosis; therapy; transgenic animal; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
CDS 366..1715
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FT mutation 787
FT /*tag= p
FT /*note= "A to T FAD mutation site (Asn141Ile)"
FT mutation 1080
FT /*tag= q
FT /*note= "A to G FAD mutation site (Met239Val)"
FT mutation 1624
FT /*tag= r
FT /*note= "T to C FAD mutation site (Ile420Thr)"

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1616 CATCTCCATCAGCTGGGCTCATCTTTTACTTCTCCACGACACACCTGG 1665
 434 alArgProPheMetAspThrLeuAlaSerHisGlnLeuTyrIle 448
 1666 TGGCGCGCTTCATGGACACCTGGCCCTCCCATCAGCTCTACATC 1709

seq_name: N_Geneseq_36.T87426

seq_documentation_block:

ID T87426 standard; DNA; 2276 BP.
 AC T87426;
 DT 07-DEC-1997 (first entry)
 DE Full AD4/AD3LP sequence.
 KW AD3: AD4/AD3LP: Alzheimer's disease; chromosome; missegregation;
 KW presentin; inhibitor; AD; trisomy 21; ss.
 OS Homo sapiens.
 PN WO9707213-A2.
 PD 27-FEB-1997.
 PF 15-AUG-1996; U13314.
 PR 16-AUG-1995; US-002448.
 PA (HARD) HARVARD COLLEGE.
 PI Li J, Potter H;
 DR P-PSDB; W28508.
 PT Identifying genes which cause chromosome missegregation - useful for
 PT identifying causes of and treatments for diseases, e.g. Alzheimer's
 PT disease, cancer and ageing
 PS Claim 28; Fig 28; 7pp; English.
 CC Identifying genes which cause improper chromosome segregation,
 CC screening for inhibitors of chromosome missegregation and processes
 CC caused by genes encoding chromosome missegregation promoters
 CC was exemplified using Alzheimer's disease. The sequences
 CC given in T87401 to T87426 can be used in the above methods.
 CC It is not clear from the figure legend, the figure and the
 CC disclosure of the specification which sequence of Fig 1 and Fig 28
 CC is the AD4/AD3LP or the AD3 sequence.
 SQ Sequence 2276 BP; 494 A; 595 C; 662 G; 525 T;

alignment_scores:

Quality: 2320.50 Length: 448
 Ratio: 5.191 Gaps: 1
 Percent Similarity: 99.777 Percent Identity: 99.777

alignment_block:

us-08-509-359B-137 x T87426 ..

Align seg 1/1 to: T87426 from: 1 to: 2276

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 411 ATGCTCATATTGCGCTCTGACAGGAGGAAGTGTGATGAGCG 460
 17 qThrSerLeuMetSerAlaGluSerProThrProArgSerCysGlnGlu 34
 461 GAGCTCCCTAATGTGCGCCGAGAGCCCCACGCGCGCTCTGCGAGGAGG 510
 34 lyArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50
 511 GCAGGCAGGCCAGAGATGGAGAGACACTGCCAGTGGAGAGCCAG 560
 51 GluAsnGluGluAspGlyGluGluAspProAspArgTyrValCysSerGl 67
 561 GAGAACGAGGAGCGGTGAGGAGGCCCTGACCGCTATGCTCTAGTGG 610
 67 yValProGlyArgProGlyLeuGluGluLeuThrLeuLysTyrG 84
 611 GGTTCGCGCGCGCCGAGCGCTGGAGAGAGAGCTGACCCCTCAAATACG 660
 84 lyAlaLysHisValIleMetLeuPheValProValThrLeuCysMetIle 100
 661 GAGCGAAGCAGCGATCATGCTGTTGTGCGCTGTCACCTCTGTGTCATGATC 710

101 ValValValAlaThrIleLysSerValArgPheTyrThrGluLysAsnGl 117
 711 GTGGTGTAGCCACCATCAAGTCTGTGGCTTCTACACAGAGAAATGG 760
 117 yGlnLeuIleTyrThrProPheThrGluAspThrProSerValGlyGlnA 134
 761 ACAGCTCATCTACAGCCATTCCTACTGAGGACACACCTCGGTGGGCCAGC 810
 134 rgLeuLeuAsnSerValLeuAsnThrLeuIleMetIleSerValIleVal 150
 811 GCCTCCTCAACTCGGTGCTGAACACCTCATCATGATGATGATGATGATG 860
 151 ValMetThrIlePheLeuValValLeuTyrLysTyrArgCysTyrLysPh 167
 861 GTTATGACCATCTTCTTGGTGGTCTCTACAAGTACCGCTGCTACAAGTT 910
 167 eileHisGlyTrpLeuIleMetSerSerLeuMetLeuLeuPheLeuPheT 184
 911 CATCATGGCTGGTGTGATCATGCTTCTCACTGATGCTGCTGCTCTCTTCA 960
 184 hrTyrIleTyrLeuGlyGluValLeuLysThrTyrAsnValAlaMetAsp 200
 961 CCTATATCTACCTTGGGGAAGTCTCAAGACCTACAATGTGGCCATGGAC 1010
 201 TyrProThrLeuLeuLeuThrValTrpAsnPheGlyAlaValGlyMetVa 217
 1011 TACCCCCCTCTTCTGCTGCTGCTGGAACCTTCGGGCGAGTGGGCATGGT 1060
 217 lCysIleHisTrpLysGlyProLeuValLeuGlnGlnAlaTyrLeuIleM 234
 1061 GTGCATCCACTGGAGGGCCCTCTGGTGTGCTGCAGCAGCCCTACCTCATCA 1110
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 301 ThrValGlyMetAlaLysLeuAspProSerSerGlnGlyAlaLeuGlnLe 317
 1311 ACGTTGGCATGGCAAGCTGGAGCCCTCTCTCAGGGTGGCCCTCCAGCT 1360
 317 uProTyrAspProGluMetGluGluAspSerTyrAspSerPheGlyGluP 334
 1361 CCCCTACGACCCGGAGATG...GAAGACTCCTATGACAGTGTGGGGAGC 1407
 334 roSerTyrProGluValPheGluProProLeuThrGlyTyrProGlyGlu 350
 1408 CTTTCATACCCGAGAGTCTTTGAGGCTCCCTTGACTGGCTACCCAGGGGAG 1457
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 1558 ACTGGAATACCACCTGGCCCTGCTTGTGGCCCATCTCTATTGGCTTGTGT 1607


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75 .LeuGluGluGluLeuThrLeuLysTyrGlyAlaLysHisValileMetL 91
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91 euPheValProValThrLeuCysMetileValValValAlaThrileLys 107
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502 TCTTTGTCTCTGACTCTGTCATGATGGTGGTGGTGGTGGTGGTGGT 551
108 SerValArgPheTyrThrGluLysAsnGlyGlnLeuIleTyrThrProPh 124
   ::::::::::::::::::::::::::::::::::::::::::::::::::::::
552 TCAGTCAGCTTTATACCGGAGGAGTGGCAGCTAATCTATACCCCAT 601
124 eThrGluAspThrProSerValGlyGlnArgLeuLeuAsnSerValLeuA 141
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602 CACAGAATACCGAGACTGTGGCCAGAGAGCCCTGCACCTCAATCTGA 651
141 snThrLeuIleMetileSerValIleValValMetThrIlePheLeuVal 157
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652 ATGCTGCCATCATGATCAGTGTGCTGATGTTGTGTCATGACTATCCCTCGTGGTG 701
158 ValLeuTyrLysTyrArgCysTyrLysPheIleHisGlyTrpLeuIleMe 174
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702 GTCTGTATAAATACAGGTGCTATAGGTCATCCATGCTGCTGCTATTAT 751
174 tSerSerLeuMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluV 191
   ::::::::::::::::::::::::::::::::::::::::::::::::::::::
752 ATCATCTCTATTGTTGCTGCTCTTTTTCATTCATTCATTCATTCATTC 801
191 aLeuLysThrTyrAsnValAlaMetAspTyrProThrLeuLeuLeuThr 207
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802 TCTTTAAACCTATAACGTTGCTGTGGACTACATATTACTGTGTCACCTCGT 851
208 ValTyrAsnPhelGlyAlaValGlyMetValCysIleHisTrpLysGlyPr 224
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852 ATCTGGAATTTGGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 901
224 oLeuValLeuGlnGlnAlaTyrLeuIleMetileSerAlaLeuMetAlaL 241
   ::::::::::::::::::::::::::::::::::::::::::::::::::::::
902 ACTTCGATCCAGCAGGCATATCTCATATGATGATGATGATGATGATGAT 951
241 euValPheIleLysTyrLeuProGluTrpSerAlaTrpValIleLeuGly 257
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952 TGGTGTATTATCAAGTACCTCCCTGATGAGTGCCTGGTGCATCTCTGGCT 1001
258 AlaIleSerValTyrAspLeuValAlaValLeuCysProLysGlyProLe 274
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1002 GTGATTTTCATGATATGATTTAGTGGTGGTGGTGGTGGTGGTGGTGGT 1051
274 uArgMetLeuValGluThrAlaGlnGluArgAsnGluProIlePheProA 291
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1052 TCGTATGCTGGTGAACAGCTCAGAGAGAAATGAACGCTTTTCCAG 1101
291 laLeuIleTyrSerSerAlaMetValTrpThrValGlyMetAlaLysLeu 307
   ::::::::::::::::::::::::::::::::::::::::::::::::::::::
1102 CTCTCATTTACTCTCAACATGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 1151
308 AspProSerSerGlnGlyAlaLeu.....GlnLeuProTyrAspProGl 322
   ::::::::::::::::::::::::::::::::::::::::::::::::::::::
1152 GACCGGGAAGCTCAAGGAGAGTATCCAAAAATCCAAAGTATATGACAGA 1201
322 u...MetGluGluAspSerTyrAspSerPheGlyGlu.....PROS 335
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335 exTyrProGluValPheGluProProLeuThrGlyTyrProGly..... 349
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1252 GATTTCAGTGAAGTGGGAGCCAGGAGGACAGTCTATCTAGGCTCAT 1301
350 .....GluGluLeu..... 352
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seq_name: N_Geneseq_36:V17358

seq_documentation_block:

ID V17358 standard; DNA; 2764 BP.

AC V17358;

DT 04-JUN-1998 (first entry)

DE PS1/467 protein coding sequence.

KW Presentin peptide; PS1/429; immunogen; immune response; PS1 gene;

KW Alzheimer's disease; mitochondrial pathology; neurodegeneration;

KW apoptosis; PS1/467; ss.

OS Homo sapiens.

FH Key Location/Qualifiers

FT CDS 249..1652

PN /*tag= a

PD W09746678-Al.

PF 11-DEC-1997.

PR 03-JUN-1997; U09272.

PR 18-JUL-1996; US-683315.

PR 06-JUN-1996; US-659296.

PA (FARB) BAYER CORP.

PI Chisholm JC, Davis JM, Drache B;

DR WPI: 98-042186/04.

DR P-PSDB; W41430.

PT DNA encoding presentin peptide PS1/429 and its analogues - useful

PS for diagnosis and treatment of Alzheimer's disease

PS Disclosure; Fig 2; 77pp; English.

CC This sequence encodes the PS1/467 presentin peptide. This sequence is

CC specifically stated as not being in the nucleic acid of the invention,

CC which encodes the PS1/429 presentin peptide PS1/429 (ii). Cells

CC transformed with the DNA are used to produce recombinant (ii) and

CC analogues, useful e.g. as immunogens for generating an immune response

CC against PS1/429. (ii) is a new product of the PS1 gene, mutations in

CC which cause Alzheimer's disease (AD). The nucleic acids are generally

CC useful as probes for detection and quantification of PS1/429,

CC particularly for diagnosis of AD, especially the target sequences that

CC hybridize with probes are isolated for sequencing. Antibodies (Ab) can

CC also be diagnosed at the protein level using Ab as immunoassay reagents.

CC Ab can also be used to identify epitopes and for affinity purification of

CC peptides. Antisense nucleic acid may also be used to regulate expression

CC of the PS1/429 gene, and both nucleic acids and peptides are useful as

CC size markers in electrophoresis, chromatography etc. The transgenic

CC animals are used as models for AD, e.g. for testing drugs. Regulators of

CC the PS1/429 gene or polypeptide can be used to treat e.g. AD or diseases

CC involving mitochondrial pathology, apoptosis and neurodegeneration.

CC Typical regulators are antisense sequences, ribozymes, aptamers,

CC synthetic or natural compounds. (ii) may also be used to target other

CC coding sequences to particular cellular locations.

SQ Sequence 2764 BP; 715 A; 624 C; 653 G; 772 T;


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FT FT      /*tag= e
FT FT      /number= 5
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FT FT      /*tag= g
FT FT      /number= 7
FT FT      797. .1017
FT FT      /*tag= h
FT FT      /number= 8
FT FT      1018. .1116
FT FT      /*tag= i
FT FT      /number= 9
FT FT      1117. .1203
FT FT      /*tag= j
FT FT      /number= 10
FT FT      1204. .1377
FT FT      /*tag= k
FT FT      /number= 11
FT FT      1378. .1496
FT FT      /*tag= l
FT FT      /number= 12
FT FT      1497. .2765
FT FT      /*tag= m
FT FT      /number= 13
FT FT      324. .335
FT FT      /*tag= n
FT FT      /note= "deletion of 12 nucleotides from 3' end of
FT FT      exon 4 by alternative splicing, deletes
FT FT      Val26-Gln-29 (not critical to PSI
FT FT      function)"
FT FT      1018. .1116
FT FT      /*tag= o
FT FT      /note= "absence of exon 9 through splicing
FT FT      variation, results in Asp-257 changing
FT FT      to Ala and fusion of Ala-257 to Thr-291"
FT FT      492
FT FT      /*tag= p
FT FT      /note= "G to C FAD mutation site (Val82Leu)"
FT FT      591
FT FT      /*tag= q
FT FT      /note= "T to C FAD mutation site (Tyr115His)"
FT FT      664
FT FT      /*tag= r
FT FT      /note= "T to C FAD mutation site (Met139Thr)"
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FT FT      /*tag= s
FT FT      /note= "T to C FAD mutation site (Ile143Thr)"
FT FT      684
FT FT      /*tag= t
FT FT      /note= "A to C FAD mutation site (Met146Leu)"
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FT FT      /*tag= z
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FT FT      /note= "C to G FAD mutation site (Leu286Val)"

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FT mutation 1399
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FT FT      /note= "G to C FAD mutation site (Gly384Ala)"
FT mutation 1422
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FT FT      /note= "C to G FAD mutation site (Leu392Val)"
FT mutation 1477
FT FT      /*tag= ad
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FT mutation 1563
FT FT      /*tag= ae
FT FT      /note= "A to G FAD mutation site (Ile439Val)"
FT WO9801549-A2.
FT PN 15-JAN-1998.
FT PD 04-JUL-1997; CA0475.
FT PF 02-JAN-1997; US-034590.
FT PR 05-JUL-1996; US-021673.
FT PR 12-JUL-1996; US-021700.
FT PR 08-NOV-1996; US-029895.
FT PA (HSCR-) HSC RES & DEV LP.
FT PA (UTOR ) UNIV TORONTO GOVERNING COUNCIL.
FT PI Fraser PE, Rommens JM, St George-Hyslop PH;
FT DR P-PSDB; W23964.
FT DR P-PSDB; W23964.
FT PT New isolated mutant presenilin-1 genes - useful for developing
FT PT products for use in detection, diagnosis and therapy of Alzheimer's
FT PT disease and for drug screening
FT PS Disclosure; Page 178-180; 238pp; English.
FT CC This cDNA clone, deposited as ATCC 97124, codes for human
FT CC presenilin-1 (hpsl, see W23964). Mutations in the presenilin genes
FT CC have been linked to the development in humans of forms of familial
FT CC Alzheimer's disease (FAD) and may be causative of other disorders,
FT CC e.g. cognitive, intellectual, neurological or physiological
FT CC disorders such as cerebral haemorrhage, schizophrenia, depression,
FT CC mental retardation and epilepsy. Isolation of the hpsl cDNA
FT CC of a physical contig spanning the AD3 region at 14q24.3, construction
FT CC and analysis of candidate genes, and recovery of candidate genes by
FT CC RT-PCR from brain mRNA. In this cDNA, exon 1 is spliced directly
FT CC to exon 3. Other hpsl sequences (see also V04667) result from
FT CC alternative splicing of the mRNA transcript. A mouse PSI homologue
FT CC (see V04668) and a human presenilin-2 sequence (see V04669) are
FT CC also provided, as well as genomic sequences for hpsl (see
FT CC T99861-71). Use of the nucleic acids and proteins comprising or
FT CC derived from the presenilins is made in screening and diagnosing
FT CC FAD, identifying and developing therapeutics for treatment of FAD,
FT CC and in producing cell lines and transgenic animals useful as models
FT CC of FAD. Nucleic acids (see V04674-80) encoding presenilin
FT CC interacting proteins are also provided.
FT SQ Sequence 2765 BP; 715 A; 624 C; 652 G; 773 T;

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alignment_scores:

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Quality: 1467.00      Length: 466
Ratio: 3.954          Gaps: 8
Percent Similarity: 79.614      Percent Identity: 65.665

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alignment_block:

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US-08-509-359B-137 x V04666

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Align seg 1/1 to: V04666 from: 1 to: 2765

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24 GluSerProThrProArgSerCysGlnGluArgGlnGlyProGluAs 40
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255 GAGTACCTGCACCGTTGTCTACTTCCAGAAATGCACAGATGCTGAGGA 304
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
40 pGlyGluAsnThrAlaGlnTrpArgSerGlnGluAsnGluGluAspGlyG 57
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
305 CAACACCTGAGCAATACTGTACTGTCAGCAATGACAATAGAGAAGCGC 354
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
57 LuGluAspProAspArgTyrValCysSerGlyValPro..... 69
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
355 AGGAGCACAACGACAGACGG...AGCCTTGCCACCTGAGCCATTATCT 401

```


CC Antibodies against the mutant polypeptide can also be used for this
 CC purpose. Vectors containing or expressing a nucleic acid molecule,
 CC protein or antibody specific for mutant P51 can be administered to a
 CC patient to reduce the likelihood, or delay the onset, of Alzheimer's
 CC disease, e.g. anti-sense RNA expression can be used to decrease
 CC expression of the P51 peptide. Transgenic animals expressing the
 CC Alzheimer's disease protein can be used to test candidate therapeutics
 CC and to investigate the normal role of P51. The P51 peptide may also be
 CC included in pharmaceutical compositions (vaccines) for Alzheimer's
 CC disease therapy.
 SQ Sequence 2765 BP; 715 A; 625 C; 652 G; 772 T;

alignment_scores:
 Quality: 1462.00 Length: 466
 Ratio: 3.941 Gaps: 8
 Percent Similarity: 79.614 Percent Identity: 65.451

alignment_block:

US-08-509-359B-137 x T85332 ..

Align seg 1/1 to: T85332 from: 1 to: 2765

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255 GAGTACCTGCACCGTGTCTCTACTTCCAGATGCACAGATGTCGAGGA 304
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
40  pGlyGluAsnThrAlaGlnTrpArgSerGlnGluAsnGluGluAspGly 57
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
305 CAACCACTGAGCAATACTGTACCTAGCAGATGACAAATAGAGAACGGC 354
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
57  luGluAspProAspArgTyrValCysSerGlyValPro..... 69
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
355 AGGAGCACACGACAGACGG...AGCCTGGCCACCTGAGCCATTATCT 401
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
70  ...GlyArgProGly..... 74
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
402 AATGGACGACCCAGGTAACTCCCGGCGAGTGGTGAGCAAGATGAGGA 451
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
75  .LeuGluGluLeuThrLeuLysTyrGlyAlaLysHisValIleMetL 91
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
452 AGAAGATGAGGAGCTGACATTGAAATATGGCGCAAGCATGATGATGC 501
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
91  euPheValProValThrLeuCysMetIleValValAlaThrIleLys 107
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
502 TCTTTGTCCCTGTGACTCTCTGATGTTGGTGGTGGTCTGCTGCTAAC 551
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
108 SerValArgPheTyrThrGluLysAsnGlyGlnLeuIleTyrThrPro 124
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
552 TCAGTCAGCTTTTATACCCGGAAGATGGGCGAGCTAATCTATACCCCA 601
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124 eThrGluAspThrProSerValGlyGlnArgLeuLeuAsnSerValLeu 141
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602 CACAGAAGATACCGAGACTGTGGCCGAGAGACCTGCACCTCAATTCGA 651
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141 snrhrLeuIleMetIleSerValIleValValMetThrIlePheLeuVal 157
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
652 ATGCTGCGCATCATGATCAGTGTCTATTGTCATGACTATCCTCCTGCTG 701
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158 ValLeuTyrLysTyrArgCysTyrLysPheIleHisGlyTyrPheLeu 174
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
702 GTTCTGTAATAACAGGTGTATAGGTATCCATGCTGCTGCTGCTATAT 751
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208 ValTrpAsnPheGlyAlaValGlyMetValCysIleHisTrpLysGlyPr 224
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852 ATCTGGAATTTTGGTGTGGTGAATGATTTCATTCACCTGCTGCTGCTGCT 901
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224 oLeuValLeuGlnGlnAlaTyrLeuIleMetIleSerAlaLeuMetAla 241
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902 ACTTCGACTCCAGGAGCATATCTCATTATGATTGATGATGATGATGATG 951
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241 euValPheIleLysTyrLeuProGluTyrPheSerAlaTyrPheValIle 257
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952 TGGTGTATTCAAGTACCTCCCTGATGAGTGGTGGTGGTGGTGGTGGT 1001
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258 AlaIleSerValTyrAspLeuValAlaValLeuCysProLysGlyProLe 274
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308 AspProSerSerGlnGlyAlaLeu.....GlnLeuProTyrAspProG 322
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335 erTyrProGluValPheGluProProLeuThrGlyTyrProGly..... 349
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353 .....GluGluGluGluGluArgGlyValLysLeuGlyLeuGlyA 366
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366 spPheIlePheTyrSerValLeuValGlyLysAlaAlaThrGlySer 382
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399 uCysLeuThrLeuLeuLeuAlaValPheLysLysAlaLeuProAlaL 416
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1502 GTGCTTACATTAATTAATTCCTTCCATTTTCAAGAAGCATTTGCCAGCTC 1551
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416 euProIleSerIleThrPheGlyLeuIlePheTyrPheSerThrAspAsn 432
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1552 TTCCAATCTCCATCACCTTGGGCTGTTTCTACTTTGCCACAGATTAT 1601
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433 LeuValArgProPheMetAspThrLeuAlaSerHisGlnLeuTyrIle 448
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1602 CTTGTACAGCCCTTTTATGGACCAATTAGCATTCATCAATTTTATATC 1649
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seq_name: N_Geneseq_36:V29525

seq_documentation_block:

ID V29525 standard; cDNA; 2764 BP.

AC V29525;

DT 13-Oct-1998 (first entry)

DE Homo sapiens PS-1 cDNA.

KW PS-1; presenilin; presenilin-1; PSP-1; Alzheimer's disease;

OS serine protease; neurodegeneration; predisposition; diagnosis; ss.

FH Homo sapiens.

Location/Qualifiers

seq_documentation_block:
 ID T59535 standard; CDNA; 1762 BP.
 AC T59535;
 DT 07-MAY-1997 (first entry)
 DE Human early onset Alzheimer's disease (EOAD) gene.
 KW Early onset Alzheimer's disease; EOAD; neurodegenerative disease;
 OS diagnosis; gene therapy; antisense; ds.
 FH Homo sapiens.
 FT Key Location/Qualifiers
 FT cds 174..1577
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 FT /label= VRSQ
 FT /note= "splice variant"
 PN WO9703086-Al.
 PD 30-JAN-1997.
 PF 26-JUN-1996; U11064.
 PR 13-JUL-1995; US-001142.
 PR 18-JUL-1995; US-001501.
 PA (USF-) UNIV SOUTH FLORIDA.
 PI Hardy JA;
 DR WPI; 97-118980/11.
 DR P-PSDB; W11839.
 PT Early onset Alzheimer's disease gene - useful for diagnosing a
 PT pre-disposition to Alzheimer's disease
 PS Claim 1; Fig 1; 44pp; English.
 CC A full-length cDNA (T59535) of the early onset Alzheimer's disease
 CC (EOAD) gene sequence codes for a 467-amino acid polypeptide (W11839).
 CC Another full-length cDNA (T59536) of an EOAD splice variant gene
 CC codes for a 463-amino acid polypeptide (W11840). The 2 sequences
 CC can be used to generate primers and probes for the diagnosis of
 CC pre-disposition to Alzheimer's disease, esp. EOAD. They can also be
 CC used for prodn. of EOAD polypeptides in transformed host cells, and
 CC antisense sequences can be used for the treatment of EOAD.
 SQ Sequence 1762 BP; 442 A; 389 C; 430 G; 478 T;

alignment_scores:
 Quality: 1454.00 Length: 466
 Ratio: 3.919 Gaps: 9
 Percent Similarity: 79.614 Percent Identity: 65.451

alignment_block:
 US-08-509-359B-137 x T59535 ..
 Align seg 1/1 to: T59535 from: 1 to: 1762

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40  pGly.....GluAsnThrAlaGlnTrpArgSerGlnGluAsnGluGlu 55
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55  spGlyGluGluAspProAspArgTyrrValCysSerGlyValPro..... 69
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277  GGCAGGAGCACACGACAGACGCGCTTGGCCACCTGAGCCATTATCT 326
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327  AATGAGACGCCCGGCTACTCCCGCAGGTGGTGAGCAAGATGAGGA 376
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377  AGAAGATGAGGAGCTGACATTGAATATGCGCCCAAGCATGTGATCATGC 426
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91  euPheValProValThrLeuCysMetIleValValValAlaAlaThrIleLys 107
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427  TCATTGTCCTGTGACTCTCTGCATGGTGGTGGTGGCTACCATTAAG 476
  
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108  SerValArgPheTyrrThrGluLysAsnGlyGlnLeuIleTyrrThrProPh 124
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477  TCAGTCAGCTTTTATACCCGGAAGATGGGAGCTAATCTATATACCCCAT 526
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124  eThrGluAspThrProSerValGlyGlnArgLeuLeuAsnSerValLeuA 141
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527  CACAGAAGATACCAGACTGTGGCCAGAGAGCCCTGCACATCAATTCGA 576
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141  snThrLeuIleMetIleSerValIleValMetThrIlePheLeuVal 157
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
577  ATGTCGCCATCATGATCAGTGTCTATTGTTGTCATGACTATCTCTCTGGT 626
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158  ValLeuTyrrLysTyrrArgCysTyrrLysPheIleHisGlyTrpLeuIle 174
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627  GTTCTCTATAAATACAGTGTCTAAGTCTATCCATGCTGCTGGCTATTAT 676
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174  tSerSerLeuMetLeuLeuPheLeuPheThrTyrrIleTyrrLeuGlyGlu 191
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368 lePheTyrSerValLeuValGlyLysAlaAlaAlaThrGlySerGlyAsp 384
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401 uThrLeuLeuLeuAlaValPheLysLysAlaLeuProAlaLeuProI 418
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167 eIleHisGlyTrpLeuIleMetSerSerLeuMetLeuLeuPheLeuPheT 184
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234 eIleSerAlaLeuMetAlaLeuValPheIleLysTyrLeuProGluTrp 250
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267 lLeuCysProLysGlyProLeuArgMetLeuValGluThrAlaGlnGluA 284
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; Sequence 28, Application US/08875972
; Patent No. 5985564
; GENERAL INFORMATION:
; APPLICANT: Huntington Potter and Jinhue Li
; TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02173-4799
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/875,972
; FILING DATE: 08-AUG-97
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/002,448
; FILING DATE: 16-AUG-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan Esq., Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: HU95-03PA
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781) 861-6240

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; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/875,972
; FILING DATE: 08-AUG-97
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/002,448
; FILING DATE: 16-AUG-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan Esq., Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: HU95-03PA
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781) 861-6240
; TELEFAX: (781) 861-9540
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1417 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: Double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 2..1129
; US-08-875-972-1

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seq_name: /cgn_6/ptodata/2/ina/5C_COMB.seq:US-08-967-101-135
seq_documentation_block:
; Sequence 135, Application US/08967101
; Patent No. 5840540
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston

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1486 CCTCCCATCTCCATCATCCTCGGGCTCGTGTCTTACTTCGCCACGGAT 1535
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seq_documentation_block:
; Sequence 135, Application US/08592541
; Patent No. 5986054
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSES: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
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; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 135:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1962 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-592-541-135

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Ratio: 3.915 Gaps: 6
Percent Similarity: 80.300 Percent Identity: 64.026

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: MEDIUM TYPE: Floppy disk
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: OPERATING SYSTEM: PC-DOS/MS-DOS
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: SOFTWARE: Patentin Release #1.0, Version #1.30
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: CURRENT APPLICATION DATA:
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: APPLICATION NUMBER: US/08/592,541
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: FILING DATE:
:
: CLASSIFICATION: 800
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: ATTORNEY/AGENT INFORMATION:
:
: NAME: Pitcher, Edmund R.
:
: TELECOMMUNICATION INFORMATION:
:
: TELEPHONE: (617) 248-7000
:
: TELEFAX: (617) 248-7100
:
: INFORMATION FOR SEQ ID NO: 133:
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: SEQUENCE CHARACTERISTICS:
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: LENGTH: 2791 base pairs
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: TYPE: nucleic acid
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: STRANDEDNESS: single
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: MOLECULE TYPE: CDNA
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seq_documentation_block:
; Sequence 1, Application US/08592541
; Patent No. 5986054
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2791 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-592-541-1

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Ratio: 3.938 Gaps: 8
Percent Similarity: 79.614 Percent Identity: 65.451

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; Sequence 3, Application US/08670964
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GENERAL INFORMATION:

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Date: Mar 18, 2000 11:52 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-1998 Compugen Ltd.

Command line parameters:

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Search information block:

Query: US-08-509-359B-137

Query length: 448

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Database sequences: 4538634

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Search time (sec): 350.460000

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ACCESSION AI925372

VERSION AI925372.1 GI:5661336

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (bases 1 to 703)

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

On May 18, 1998 this sequence version replaced gi:3137011.

Contact: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert.Strausberg@nih.gov

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

DNA Sequencing by: Greg Lennon, Ph.D.

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.lnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco

High quality sequence used: 459.

Location/Qualifiers

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differentiated (4 pooled tumors, including primary and

metastatic)"

/dev_stage="adult"

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/note="Organ: lung; Vector: pT73D-Pac (Pharmacia) with a

modified polylinker; 1st strand cDNA was prepared from

pooled lung tumor tissue, and was then primed with a Not I

- oligo(dT) primer. Double-stranded cDNA was ligated to

Eco RI adaptors (Pharmacia), digested with Not I and

cloned into the Not I and Eco RI sites of the modified

pT73 vector. Library went through one round of

normalization. Library constructed by Bento Soares and M.

Fatima Bonaldo."

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ORIGIN

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Ratio: 5.088 Gaps: 0

Percent Similarity: 97.425 Percent Identity: 97.425

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US-08-509-359B-137 x AI925372

Align seg 1/1 to: AI925372 from: 1 to: 703

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452 CAGGCTACCTCATCATGATGATGCGCTCATGGCCCTAGTGTTCATCAA 501

245 sTyrLeuProGluTtpSerAlaTtpValIleLeuGlyAlaIleSerValT 262

502 GTACCTCCAGAGTGGTCCGGTGGGTGATCTCGGGCCATCTCTGTGT 551

262 yAspLeuValAlaValLeuCysProLysGlyProLeuArgMetLeuVal 278

552 ATGATCTCGTGGCTGTGTGTGCCANAGGCCCTCTGAGAATGCTGTA 601

279 GluThrAlaGlnGluArgAsnGluProIlePheProAlaLeuIleTyrSe 295

602 GAAACTGCCAGGAGAGAAATGAGCCCATATTCCCTGCTTCATATACTC 651

295 rSerAlaMet 298

652 ATCTGNCATG 661

seq_name: gb_est35:AI831581

seq_documentation_block:

LOCUS AI831581 680 bp mRNA EST 26-AUG-1999
DEFINITION wj39q04.x1 NCI-CGAP_Lu19 Homo sapiens cDNA clone IMAGE:2405191 3',
similar to SW:PSN2_HUMAN P49810 PRESENILIN 2 ;, mRNA sequence.
ACCESSION AI831581
VERSION AI831581.1 GI:5452252
KEYWORDS EST.
SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 680)
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

JOURNAL

Unpublished (1997)

On Dec 20, 1995 this sequence version replaced gi:1130797.

Contact: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert.Strausberg@nih.gov

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco

High quality sequence stop: 456.

FEATURES

source

1. .680

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:2405191"

/clone_lib="NCI-CGAP_Lu19"

/tissue_type="squamous cell carcinoma, poorly

differentiated (4 pooled tumors, including primary and

metastatic)"

/dev_stage="adult"

/lab_host="DH10B (phage-resistant)"

/note="Organ: lung; Vector: p7T3D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from pooled lung tumor tissue, and was then primed with a Not I - oligo(drf) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified p7T3 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 137 a 191 c 173 g 178 t 1 others
ORIGIN

alignment_scores:

Quality: 1118.00 Length: 226
Ratio: 5.059 Gaps: 0
Percent Similarity: 97.788 Percent Identity: 96.903

alignment_block:

US-08-509-359B-137 x AI831581

Align seg 1/1 to: AI831581 from: 1 to: 680

79 LeuThrLeuLysTyrGlyAlaLysHisValIleMetLeuPheValProVa 95
|||||
3 CTGACCCCTCAAACTACTGATCGAGCAGCGATCATGCTGTTGTGCGCTGT 52
95 lThrLeuCysMetIleValValAlaThrIleLysSerValArgPheT 112
|||||
53 CACTCTGTCATGATCGTGGTAGCCACCACCAAGTCTGTGCGCTTCT 102
112 yThrGluLysAsnGlyGlnLeuIleTyrThrProPheThrGluAspThr 128
|||||
103 ACACAGAGAAGAAATGACAGCTCATCTACAGCCATCTACTGAGGACACA 152
129 ProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLeuIleMe 145
|||||
153 CCCTCGTGGGCCAGGCGCTCTCAACTCCGTGCTGAACACCCCTCATCAT 202
145 tileSerValIleValValMetThrIlePheLeuValValLeuTyrLysT 162
|||||
203 GATCAGCGTCATCGTGGTATGACCATCTCTTGGTGGTCTCTACAAGT 252
162 yArgCysTyrLysPheIleHisGlyTrpLeuIleMetSerSerLeuMet 178
|||||
253 ACCGCTGCTACAAAGTTCATCCATGGCTGGTGTGATCATGCTTCTCACATG 302
179 LeuLeuPheLeuPheThrIleTyrLeuGlyGluValLeuLysThrTy 195
|||||
303 CTGCTGTTCCTCTTCCACCTATATCTACCTGGGGAAGTGTCTCAAGACCTA 352
195 rAsnValAlaMetAspTyrProThrLeuLeuLeuThrValTtpAsnPhg 212
|||||
353 CAATGGCCATGGACTACCCACCCTCTGTGCTGCTGGAACCTTCG 402
212 lyAlaValGlyMetValCysIleHisTrpLysGlyProLeuValLeuGln 228
|||||
403 GGGCAGTGGGATGTGTGATCCACTGGAAGGCCCTCTGGTGTGCAG 452
229 GlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuValPheIleLy 245
|||||
453 CAGGCTACCTCATCATGATGATGCGCTCATGGCCCTAGTGTTCATCAA 502
245 sTyrLeuProGluTtpSerAlaTtpValIleLeuGlyAlaIleSerValT 262
|||||
503 GTACCTCCAGAGTGGTCCGGTGGGTGATCTGCGCCATCTCTGTGT 552
262 yAspLeuValAlaValLeuCysProLysGlyProLeuArgMetLeuVal 278
|||||
553 ATGATCTCGTGGCTGTGTGTACCATAGGCGCTCTGAGAATGCTGGTA 602
279 GluThrAlaGlnGluArgAsnGluProIlePheProAlaLeuIleTyrSe 295
|||||

High quality sequence stop: 397.

FEATURES

Location/Qualifiers
1..597
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2619929"
/tissue_type="NCI_CGAP_Brn50"
/tissue_type="medulloblastoma"
/lab_host="DH10B (phage resistant)"
/note="Organ: brain; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from medulloblastoma tumor tissue, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. This library is normalized. Library constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 119 a 170 c 147 g 161 t

alignment_scores:

Quality: 916.00 Length: 199
Ratio: 4.796 Gaps: 0
Percent Similarity: 95.980 Percent Identity: 92.462

alignment_block:

US-08-509-359b-137 x AW131752

Align seg 1/1 to: AW131752 from: 1 to: 597

79 LeuThrLeuLysTyrGlyAlaLysHisValIleMetLeuPheValProva 95
|||||
2 CTGACCCCTCAATACGTG.TCGAAGCACGTGATCATGCTTTGTGCCCTGT 50
95 lThrLeuCysMetIleValValAlaThrIleLysSerValArgPheT 112
51 CACCTGTGTCATGATCGTGGTGGTAGCCACCATCAAGTCTGCGCTTCT 100
112 yThrGluLysAsnGlyGlnLeuIleTyrThrProPheThrGluAspThr 128
101 ACACAGAGAAGAAATGGACAGCTCATCTACACGCCATTCTCAGGACACA 150
129 ProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLeuIleMe 145
151 CCCTCGGTGGGCCAGCGCTCTCTCAACTCCGTGCTGAACCCCTCATCAT 200
145 tIleSerValIleValValMetThrIlePheLeuValValLeuTyrLysT 162
201 GATCAGCTCATCGTGGGTATGACCATCTTCTGTGGTGTCTCTACAAAGT 250
162 yArgCysTyrLysPheIleHisGlyTrpLeuIleMetSerSerLeuMet 178
251 ACCGCTGTACAAGTTTCATCCATGCTGGTTCATCATGCTTCATCATGATG 300
179 LeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluValLeuLysThrTy 195
301 CTGCTGTTCTCTTCCATCATATATCTACCTTGGGAAGTGTCTCAAGACCTA 350
195 rAsnValAlaMetAspTyrProThrLeuLeuThrValTrpAsnPheG 212
351 CAATGTGCCCATGGACTACCCACCCCTTCTGCTGACTGTCTGGAACCTCG 400
212 lAlaValGlyMetValCysIleHisTrpLysGlyProLeuValLeuGln 228
401 GGGCAGTGGGCATGGTGTGCATCCACTCGAAGGGCCCTCTGGTGTGCGAG 450
229 GlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuValPheIleTy 245
451 CACGCCTACCTCATCATCATGATCAGTGCCTCATGTCCTATTGTTTCATCAA 500
245 sTyrLeuProGluTrpSerAlaTrpValIleLeuGlyAlaIleSerValT 262
| |||||

501 GATCTACCAAAAGTGTACGCGTGGTGCATCTGAGCGCATATCTGTGT 550

262 yrAspLeuValAlaValLeuCysProLysGlyProLeuArgMetLeu 277

|||||
551 ATGATCTCGTCCCTGGCTGTGTACCAATGGCCTGTGTGAATGCTG 597

seq_name: gb_est17:AA602396

seq_documentation_block:

LOCUS AA602396 538 bp mRNA EST 08-OCT-1997
DEFINITION nc30d05.s1 NCI_CGAP_Pr22 Homo sapiens cDNA clone IMAGE:1102185 3'
Similar to TR:G1244640 G1244640 PRESENILIN I-374. ;, mRNA sequence.
ACCESSION AA602396
VERSION AA602396.1 GI:2436374
KEYWORDS EST.
SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS
1 (bases 1 to 538)
TITLE
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
NATIONAL Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
On Sep 19, 1997 this sequence version replaced gi:1517341.

Contact: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert.Strausberg@nih.gov

Tissue Procurement: Michael J. Brownstein, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 1338 Std Error: 0.00

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 404.

FEATURES

source

1..538
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1102185"
/clone_lib="NCI_CGAP_Pr22"
/sex="male"
/tissue_type="normal prostate"
/lab_host="DH10B"
/note="Organ: prostate; Vector: pT73D-Pac (Pharmacia)
from normal prostate bulk tissue, and was then primed with
a Not I - oligo(dT) primer. Double-stranded cDNA was
ligated to Eco RI adaptors (Pharmacia), digested with Not
I and cloned into the Not I and Eco RI sites of the
modified pT73 vector. Library is normalized, and was
constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 101 a 155 c 138 g 143 t 1 others
ORIGIN

alignment_scores:

Quality: 890.00 Length: 179
Ratio: 5.028 Gaps: 0
Percent Similarity: 98.883 Percent Identity: 98.324

alignment_block:

US-08-509-359b-137 x AA602396

Align seg 1/1 to: AA602396 from: 1 to: 538

94 ProValThrLeuCysMetIleValValAlaThrIleLysSerValAr 110

|||||

```

DEFINITION wh6e09.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:2385736 3'
similar to SW:PSN2_HUMAN P49810 PRESENIIN 2 ;, mRNA sequence.
ACCESSION A1765870
VERSION A1765870.1 GI:5232379
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 707)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT On Feb 17, 1998 this sequence version replaced gi:2889754.
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone Distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco
High quality sequence stop: 466.
Location/Qualifiers
1..707
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2385736"
/clone.lib="NCI_CGAP_Kid11"
/lab.host="DH10B"
/note="Organ: kidney; Vector: pT73D-Pac (Pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Plasmid DNA from the normalized library NCI_CGAP_Kid3 was
prepared, and ss circles were made in vitro. Following HAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(cloneids 1323376-1323911, 1456007-1456775, and
1500552-1502855). Subtraction by Bento Soares and M.
Fatima Bonaldo."
BASE COUNT 177 a 197 c 182 g 150 t 1 others
ORIGIN

alignment_scores
Quality: 870.00 Length: 186
Ratio: 4.807 Gaps: 0
Percent Similarity: 97.312 Percent Identity: 95.161

alignment_block:
US-08-509-359b-137 x A1765870/rev ..

Align seg 1/1 to reverse of: A1765870 from: 1 to: 707

263 AspLeuValAlaValLeuCysProLysGlyProLeuArgMetLeuValG1 279
||||| ||||||| ||||||| ||||||| ||||||| |||||||
707 GATCTGGGGCGTGTGCTGTCCAAAGGCGTCTTGAGATGCTGGTAGA 658

279 uThrAlaGlnGluArgAsnGluProIlePheProAlaLeuIleTyrSers 296
||||| :||||: ||||||| ||||||| ||||||| |||||||
657 ACTGCCAGAGAGAAATGAGCCCATATCTCCCTGCCCTGATATA. NCAT 609

296 exAlaMetValTrpThrValGlyMetAlaLysLeuAspProSerSerGln 312
||||| ||||||| ||||||| ||||||| ||||||| |||||||
608 CTGCCATGTTGGAGCGTGGCATGGGAAGCTGGACCCCTCC.TCTCAG 560

313 GlyAlaLeuGlnLeuProTyrAspProGluMetGluGluAspSerTyrAs 329

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||||| ||||||| ||||||| ||||||| ||||||| |||||||
559 GTGTCCTCCAGCTCCCTACGACCCGGAGATGGAGAGACTCCTATGA 510

329 pSerPheGlyGluProSerTyrProGluValPheGluProProLeuThrG 346
||||| ||||||| ||||||| ||||||| ||||||| |||||||
509 CAGTTTTGGGGAGCCTTCATACCCCGAAGCTTTTGAGCCTCCCTTGACTG 460

346 lYrProGlyGluGluLeuGluGluGluGluGluGluGluGluGluGlu 362
||||| ||||||| ||||||| ||||||| ||||||| |||||||
459 GCTACCCAGGGAGGAGCTGGAGAGAGAGAGAGAGAGAGAGAGAGAGCTT 410

363 GlyLeuGlyAspPheIlePheTyrSerValLeuValGlyLysAlaAlaAl 379
||||| ||||||| ||||||| ||||||| ||||||| |||||||
409 GGCTCTGGGAGACTTCATCTTCTACAGTGTGCTGTGGCAGAGGGCTGC 360

379 aThrGlySerGlyAspTrpAsnThrThrLeuAlaCysPheValAlaIleL 396
||||| ||||||| ||||||| ||||||| ||||||| |||||||
359 CACGGGCGAGGGGAGCTGGAATACACAGCTGGCCTTCGTGGCCATCC 310

396 euIleGlyLeuCysLeuThrLeuLeuLeuLeuAlaValPheLysLysAla 412
||||| ||||||| ||||||| ||||||| ||||||| |||||||
309 TCATTGGCTTGTGTCTGACCTCTCTGCTGTGCTGTGTTCAAGAAGCG 260

413 LeuProAlaLeuProIleSerIleThrPheGlyLeuIlePheTyrPheSe 429
||||| ||||||| ||||||| ||||||| ||||||| |||||||
259 TGCCCCGCCCTCCCAATTCATCACGTTGGGCTCATCTTTTACTTCTC 210

429 rThrAspAsnLeuValArgProPheMetAspThrLeuAlaSerHisGlnL 446
||||| ||||||| ||||||| ||||||| ||||||| |||||||
209 CACGGACAACCTGGTGGCGCGGTTCATGACACCCCTGGCCTCCCATCAGC 160

446 euTyrIle 448
|||||
159 TCTACATT 152

seq_name: gb_est44:AW212769

seq_documentation_block:
LOCUS AW212769 555 bp mRNA EST 03-DEC-1999
DEFINITION u066e03.x1 NCI_CGAP_Mam1 Mus musculus cDNA clone IMAGE:2647516 3'
similar to gb:U42177 Mus musculus S182 protein mRNA, complete cds
(MOUSE);, mRNA sequence.
ACCESSION AW212769
VERSION AW212769.1 GI:6518888
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 555)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT On Jul 7, 1999 this sequence version replaced gi:5406916.
Other_ESTs: u066e03.y1
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Washington University Genome Sequencing Center
Clone Distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

MG1:1027968
Seq primer: -40UP from Gibco
High quality sequence stop: 421.
Location/Qualifiers
1..555
/organism="Mus musculus"

```

FEATURES
source

||||| 3 CCTGTCACTCTGTGCATGTCGGTGGTAGCCACCATCAAGTCTGTGGC 52
110 gPheTyrThrGluLysAsnGlyGlnLeuLeuTyrThrProPheThrGluA 127
||||| 53 CTTCTACACAGAGAAGATGGACAGCTCATCTACACGCCATTCACCTGAGG 102
127 spThrProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLeu 143
||||| 103 ACACACCTCGTGGGCCAGCGCTCTCACTCCGCTGTGAACACCTC 152
144 IleMetIleSerValIleValValMetThrIlePheLeuValValLeuTy 160
||||| 153 ATCATCATCAGCTCATCTGGTGTATGACCATCTTCTTGGTGTGCTCTA 202
160 rLysTyrArgCysTyrLysPheIleHisGlyTrpLeuIleMetSerSerL 177
||||| 203 CAAGTACCCTGCTACAAAGTTCATCCATGGCTGGTTGATCATCTCTTCA 252
177 euMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluValLeuLys 193
||||| 253 TGAATGCTGTCTCTTCACTATATCTACCTTACCTTGGGGAAGTGTCAAG 302
194 ThrTyrAsnValAlaMetAspTyrProThrLeuLeuLeuThrValTyrP 210
||||| 303 ACTACAATGTGGCCATGACATACCCACCTCTTCTGCTGACTCTGTGAA 352
210 nPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProLeuVal 227
||||| 353 CTTGGGGGAGTGGGATGGTGTGATCCACTGGAAGGCCCTCTGGTGC 402
227 euGlnGlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuValPhe 243
||||| 403 TGACGAGGCCCTACCTCATCATGATCAGTGGCTCATGGCCCTAGTGTTC 452
244 IleLysTyrLeuProGluTyrSerAlaTyrPValIleLeuGlyAlaIle 260
||||| 453 ATCAAGTACTC.CCAGAGTGGTCCGGTGGTATC...TGGCGGCCANC 498
260 rValTyrAspLeuValAlaValLeuCysProLysGlyProLeuArgMetL 277
||||| 499 TCTGTGTATGATCCGTTNGTGTGTGTCAAAGGCT...CTGAGAATGT 545
277 euValGluThrAlaGlnGluArgAsnGluPro 287
546 GG...TAGAANTGCCAGGAGAGAATGAGCCA 574
seq_name: gb_est25:A1276606
seq_documentation_block: 481 bp mRNA EST 29-JAN-1999
LOCUS A1276606
DEFINITION gi71h07.x1 Soares_NHMPu_s1 Homo sapiens cDNA clone IMAGE:1877821
3', similar to SW:PSN2_HUMAN P49810 PRESENILIN 2 ;, mRNA sequence.
ACCESSION A1276606
VERSION A1276606.1 GI:3898880
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 481)
NCI-CCAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
On Jan 19, 1998 this sequence version replaced gi:2285413.
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Insert Length: 1769 Std Error: 0.00
Seq primer: -40UP from Gibco

High quality sequence stop: 448.
Location/Qualifiers
1..481
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1877821"
/clone_lib="Soares_NHMPu_s1"
/tissue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"
/lab_host="DH10B"
/note="Organ: mixed (see below); Vector: pT7T3D-Pac
(Pharmacia) with a modified polylinker; Site.1: Not I;
Site.2: Eco RI: Equal amounts of plasmid DNA from three
normalized libraries (melanocyte 2NHM, pregnant uterus
NDHPU, and fetal heart NDH19W) were mixed, and ss circles
were made in vitro. Following HAP purification, this DNA
was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from pools of
5,000 clones made from the same 3 libraries. The pools
consisted of I.M.A.G.E. clones 260232-265223,
340488-345479, and 484488-489479."
BASE COUNT 100 a 138 c 119 g 124 t
ORIGIN
alignment_scores:
Quality: 824.00 Length: 160
Ratio: 5.150 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000
alignment_block:
US-08-509-359B-137 x A1276606 ..
Align seg 1/1 to: A1276606 from: 1 to: 481
77 GUGluLeuThrLeuLysTyrGlyAlaLysHisValIleMetLeuPheVa 93
||||| 2 GAAGAGCTGACCTCAATACGGAGCAGCATGTGATCATGCTGTTGT 51
93 lProValThrLeuCysMetIleValValAlaThrIleLysSerValA 110
||||| 52 GCCTGTCACTCTGTGCATGATCGTGGTGGTAGCCACCATCAAGTCTGNGC 101
110 rGpheTyrThrGluLysAsnGlyGlnLeuLeuTyrThrProPheThrGlu 126
||||| 102 GCTTCTACACAGAGAGATGGACAGCTCATCTACAGCCATTCACCTGAG 151
127 AspThrProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLe 143
||||| 152 GACACACCTCGGTGGGCCAGCGCTCTCAACTCCGCTGGAACACCT 201
143 uIleMetIleSerValIleValValMetThrIlePheLeuValValLeu 160
||||| 202 CATCATGATCAGCGTCATCGTGGTATGACCATCTTCTTGGTGTGCTCT 251
160 yrlLysTyrArgCysTyrLysPheIleHisGlyTrpLeuIleMetSerSer 176
||||| 252 ACAAGTACCCTGTACAAAGTTCATCCATGGCTGGTGTGATCATGCTTCA 301
177 LeuMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluValLeuLy 193
||||| 302 CTGATGTGCTGTCTCTTCACCTATATCTACCTTGGGGAAGTGTCTCAA 351
193 sThrTyrAsnValAlaMetAspTyrProThrLeuLeuLeuThrValTyrP 210
||||| 352 GACCTCAATGTGGCCATGGACTACCCACCTCTTGTGACTGTCTGGA 401
210 snPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProLeuVal 226
||||| 402 ACTTCGGGGGAGTGGGATGGTGTGATCCACTGGAAGGCCCTCTGGTG 451
227 LeuGlnGlnAlaTyrLeuIleMetIleSer 236
|||||

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/db_xref="ATCC (inhost):86429"
/db_xref="GDB:D0S9004E"
/db_xref="taxon:9606"
/clone="IB913"
/clone_lib="Infant brain, Bento Soares"
/lab_host="E. coli DH5-alpha"
/notes="Vector: BA, M13-derived; Site_1: HindIII; Site_2:
NotI; The infant brain library, constructed by Bento
Soares, Columbia University, was oligo-(dT) primed and
directionally cloned into an M13-derived plasmid using
total brain mRNA from a 72-day old human female afflicted
with spinal muscular atrophy."
BASE COUNT 100 a 133 c 119 g 122 t 2 others
ORIGIN

alignment_scores:
  Quality: 732.50 Length: 154
  Ratio: 4.883 Gaps: 3
Percent Similarity: 97.403 Percent Identity: 96.753

alignment_block:
US-08-509-359B-137 x T03796 ..
Align seg 1/1 to: T03796 from: 1 to: 476

78 GluLeuThrLeuLysTyrGlyAlaLysHis.ValIleMetLeuPheValP 94
|||||
11 GAGCTGACCCCTCAATACGGAGCG...ANACGTNATCATGCTGTTGTGC 57
|||||
94 roValThrLeuCysMetIleValValValAlaThrIleLysSerValArg 110
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58 CTGTCACCTCTGTCATGATGCTGGTGGTAGCCACCATCAAGTCTGTGGCC 107
|||||
111 PheTyrThrGluLysAsnGlyGlnLeuIleTyrThrProPheThrGluAs 127
|||||
108 TTCACACAGAGAGAGATGGACAGCTCATCTACACGCCATTCACGTGAGGA 157
|||||
127 PThrProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLeuI 144
|||||
158 CACACCCCTCGGTGGGCCAGCGCTCTCAACTCCGCTGCTGAACCCCTCA 207
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144 LeMetIleSerValIleValValMetThrIlePheLeuValValLeuTyr 160
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208 TCATGATCAGCGTCATCGTGGTATGACCATCTCTTGGTGGTGCTCTAC 257
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161 LysTyrArgCysTyrLysPheIleHisGlyTrpLeuIleMetSerSerLe 177
|||||
258 AAGTACCCTGCTACAAAGTTATCATGCTGGTGGTTCATGCTCTTCACT 307
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177 uMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluValLeuLysT 194
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308 GATGCTGCTGTTCTCTTACCATATATCTACCTTGGGGAAGTGCTCAAGA 357
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194 hrTyrAsnValAlaMetAspTyrProThrLeuLeuLeuThrValTrp.A 210
|||||
358 CCTCAATGTGGCCATGACACTACCCACCCCTCTTCTGCTACTGTCTGGAA 407
|||||
210 nPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProLeuValL 227
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408 CTTGGGGCAGTGGGCATGGTGGTGCATCCATCTGGAAGGGGCGCTTTGGTGC 457
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227 euGlnGln 229
|||||
458 TGCAGAGG 465

seq_name: gb_est23:AI097783

seq_documentation_block:
LOCUS AI097783 483 bp mRNA EST 20-AUG-1998
DEFINITION ue35g10.y1 Sugano mouse liver mlia Mus musculus cDNA clone
IMAGE:1482402.5, similar to gb:142177 Mus musculus S182 protein
mRNA, complete cds (MOUSE);, mRNA sequence.

```

AI097783
 GI:3447308
 EST
 house mouse.
 Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 483)
 Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,
 Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,
 Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,
 Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and
 Waterston,R.
 The WashU-HHMI Mouse EST Project
 Unpublished (1996)
 On Dec 18, 1997 this sequence version replaced gi:2339814.
 Contact: Marra M/Mouse EST Project
 WashU-HHMI Mouse EST Project
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: mouseest@watson.wustl.edu
 This clone is available royalty-free through LLNL; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 MGI:930758
 Seq primer: custom primer used.
 Location/Qualifiers
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 /organism="Mus musculus"
 /strain="C57BL"
 /db_xref="taxon:10090"
 /clone="IMAGE:1482402"
 /clone_lib="Sugano mouse liver mlia"
 /sex="female"
 /dev_stage="adult"
 /lab_host="DH10B"
 /note="organ: liver; Vector: pME18S-FL3; Site_1: DraIII
 (CACTGTGTG); Site_2: DraIII (CACCATGTG); 1st strand cDNA
 was primed with an oligo(dT) primer
 [ATGTGGCTTTTCTTTTCTTTT]; double-stranded cDNA was
 ligated to a DraIII adaptor [TGTGGCTCTACTGG], digested
 and cloned into distinct DraIII sites of the pME18S-FL3
 vector (5' site CACTGTGTG, 3' site CACCATGTG). XhoI should
 be used to isolate the cDNA insert. Size selection was
 performed to exclude fragments <1.5kb. Library
 constructed by Dr. Sumio Sugano (University of Tokyo
 Institute of Medical Science). Custom primers for
 sequencing: 5' end primer CTTCGTCTCTAAAGCTGCG and 3' end
 primer CGACCTGCAGCTCGACACA."
 BASE COUNT 94 a 152 c 126 g 111 t
 ORIGIN

alignment_scores:
 Quality: 704.00 Length: 150
 Ratio: 4.993 Gaps: 0
 Percent Similarity: 94.000 Percent Identity: 90.667
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 Align seg 1/1 to: AI097783 from: 1 to: 483

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 33 GAGCATCTGTGCTCTTCTCTCTCTAGCTGCCATGTGTGGACGCT 82
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 302 lGlyMetAlaLysLeuAspProSerSerGlnGlyAlaLeuGlnLeuProf 319
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 83 GGGCATGGCAAGCTGGACCCCTCTCTCTCAAGAGCGCTGCAGCTCCCT 132
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 319 yrAspProGluMetGluGluAspSerTyrAspSerPheGlyGluProSer 335

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 20, 2000, 04:21:11 ; Search time 35.25 Seconds
(without alignments)
301.032 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFMASDSEEEVCDETSI.....STDNLVRPFMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 188963 seqs, 23686106 residues

Database : A_Geneseq_36:*

Word size : 0

Number of hits that pass the threshold : 188963

pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query		Length	DB	ID	Description
		Match					
1	2336	100.0	448	1	W05762		Human presenilin-2
2	2336	100.0	448	1	W11321		Human AD4 protein.
3	2336	100.0	448	1	W23967		Human presenilin-2
4	2332	99.8	448	1	W05763		Presenilin-2 M239V
5	2331	99.8	448	1	W05765		Presenilin-2 I420T
6	2327	99.6	448	1	W05764		Presenilin-2 N141I
7	2320.5	99.3	447	1	W28508		Full AD4/AD3LP seq
8	2136	91.4	414	1	W28506		Presenilin-2 delta
9	1623.5	69.5	376	1	W28506		AD4/AD3LP sequence
10	1468	62.8	467	1	W05735		Murine presenilin.
11	1468	62.8	467	1	W23966		Mouse presenilin-1
12	1467	62.8	467	1	W05733		presenilin-1-1. Ne
13	1467	62.8	467	1	W41430		PS1/467 protein. D
14	1467	62.8	467	1	W23964		Human presenilin-1
15	1464	62.7	467	1	W05755		Presenilin-1-1 L28
16	1464	62.7	467	1	W05758		Presenilin-1-1 L39
17	1464	62.7	467	1	W05737		Presenilin-1-1 V82
18	1464	62.7	467	1	W05746		Presenilin-1-1 I21
19	1463	62.6	467	1	W05754		Presenilin-1-1 A28
20	1463	62.6	467	1	W05736		Presenilin-1-1 A79
21	1463	62.6	467	1	W05747		Presenilin-1-1 I23
22	1463	62.6	467	1	W05749		Presenilin-1-1 A26
23	1463	62.6	467	1	W41431		Mouse PS1/467 prot
24	1462	62.6	467	1	W05738		Presenilin-1-1 V96
25	1462	62.6	467	1	W05739		Presenilin-1-1 Y11
26	1462	62.6	467	1	W05741		Presenilin-1-1 I14
27	1462	62.6	467	1	W05748		Presenilin-1-1 A24
28	1462	62.6	467	1	W27176		Human S182 gene, P
29	1461	62.5	467	1	W05753		Presenilin-1-1 E28
30	1461	62.5	467	1	W05757		Presenilin-1-1 G38
31	1461	62.5	467	1	W05740		Presenilin-1-1 M13
32	1461	62.5	467	1	W05742		Presenilin-1-1 M14
33	1460	62.5	467	1	W05744		Presenilin-1-1 L17
34	1460	62.5	467	1	W56770		Homo sapiens PS-1.
35	1459	62.5	467	1	W05752		Presenilin-1-1 P26
36	1458	62.4	467	1	W05743		Presenilin-1-1 H16
37	1458	62.4	467	1	W05745		Presenilin-1-1 G20
38	1457	62.4	467	1	W05751		Presenilin-1-1 P26
39	1456	62.3	467	1	W05759		Presenilin-1-1 C41

40 1455 62.3 467 1 W05750 Presenilin-1-1 C26
41 1454 62.2 467 1 W11839 Human early onset
42 1454 62.2 463 1 W11840 Early onset Alzheimer
43 1448 62.0 463 1 W05734 Presenilin-1-2. Ne
44 1448 62.0 463 1 W23965 Human presenilin-1
45 1446 61.9 463 1 W42375 Human presenilin I

ALIGNMENTS

RESULT 1

W05762

ID W05762 standard; Protein; 448 AA.

AC W05762;

DT 25-JUL-1997 (first entry)

DE Human presenilin-2.

KW Presenilin-2; human; hpsl-1; hpsl-2; PS-2; integral membrane protein; AD;

KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;

KW depression; antibody; gene expression modulator; therapy.

OS Homo sapiens.

PN W09634099-A2.

PD 31-OCT-1996.

PF 29-APR-1996; CA0263.

PR 28-APR-1995; US-431048.

PR 28-JUN-1995; US-496841.

PR 31-JUL-1995; US-509359.

PA (HSCR-) HSC RES & DEV LP.

PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.

PI Fraser PE, Rommens JM, St George-Hyslop PH;

DR WPI: 96-497631/49.

DR N-FSDB; T40031.

PT New presenilin genes - useful for diagnosis, therapy and drug

PT screening of familial Alzheimer's disease, cerebral disorders, etc.

PS Claim 4; Page 148-150; 178pp; English.

CC W05733 and W05734 represent the two different forms of wild type human

CC presenilin-1 (PS-1). The form represented by W05734 results from

CC alternate splicing of the genomic DNA sequence. W05735 represents the

CC coding sequence for wild type mouse PS-1. The presenilins are a family of

CC highly conserved integral membrane proteins with a common structural

CC motif, common alternate splicing patterns, and common mutational hot spot

CC regions. Mutations in PS genes are implicated in familial Alzheimer's

CC disease (AD) and possibly other diseases such as cerebral haemorrhage,

CC schizophrenia, depression etc., so detection of mutations in the DNA

CC encoding these sequences can be used for diagnosis of these diseases.

CC These proteins, or vectors that express them or containing antisense

CC sequences, antibodies selective for mutant forms of these proteins (such

CC as W05736) and modulators of PS gene expression are potentially useful

CC for treatment of AD etc. Transgenic animals are useful as models for drug

CC screening. The antibodies can also be used e.g. for affinity purification

CC and in immunoassays.

SQ Sequence 448 AA;

Query Match 100.0%; Score 2336; DB 1; Length 448;

Best Local Similarity 100.0%; Pred. No. 7.3e-237;

Matches 448; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MLTFMASDSEEEVCDETSLSMAESPTPRSCQGRGPDGENTAQRSGNEDEGEDP 60

Db 1 MLTFMASDSEEEVCDETSLSMAESPTPRSCQGRGPDGENTAQRSGNEDEGEDP 60

QY 61 DRYVCSGVPRPPGLEEEETLKYGAHVIMLFVPVTLCMIVVATIKSVRYFEKNGQLI 120

Db 61 DRYVCSGVPRPPGLEEEETLKYGAHVIMLFVPVTLCMIVVATIKSVRYFEKNGQLI 120

QY 121 YPPTEDTPSVQGRLLNSVINTLIMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMLL 180

Db 121 YPPTEDTPSVQGRLLNSVINTLIMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMLL 180

QY 181 FLFTYIIGEVKVTYNVMDYPTLLTWNFGAVGWCIIHWKGPLVLQQAYLIMISALMA 240

Db 181 FLFTYIIGEVKVTYNVMDYPTLLTWNFGAVGWCIIHWKGPLVLQQAYLIMISALMA 240

CC regions. Mutations in PS genes are implicated in familial Alzheimer's
 CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
 CC schizophrenia, depression etc., so detection of mutations in the DNA
 CC encoding the wild type sequences can be used for diagnosis of these
 CC diseases. The wild type proteins, or vectors that express them or
 CC containing antisense sequences, antibodies selective for these mutant
 CC forms of the proteins and modulators of PS gene expression are
 CC potentially useful for treatment of AD etc. Transgenic animals are useful
 CC as models for drug screening. The antibodies can also be used e.g. for
 CC affinity purification and in immunoassays.
 SQ Sequence 448 AA;

Query Match 99.6%; Score 2327; DB 1; Length 448;
 Best Local Similarity 99.8%; Pred. No. 6.4e-236;
 Matches 447; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 MLTFMADSEEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQRWSENEDEEDP 60
 Db 1 MLTFMADSEEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQRWSENEDEEDP 60

Qy 61 DRYVCSGVPGRPPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120
 Db 61 DRYVCSGVPGRPPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120

Qy 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLL 180
 Db 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLL 180

Qy 181 FLFTYIYLGELVKTYNVAMDYPTLLTWNFGAVGMVCIHWKGPLVLOQAYLIMISALMA 240
 Db 181 FLFTYIYLGELVKTYNVAMDYPTLLTWNFGAVGMVCIHWKGPLVLOQAYLIMISALMA 240

Qy 241 LVFTIKYLPENSAWVILGAISSYDVLVAVLCPKGPLRMLVETAQERNEPIFPALYSSAMVW 300
 Db 241 LVFTIKYLPENSAWVILGAISSYDVLVAVLCPKGPLRMLVETAQERNEPIFPALYSSAMVW 300

Qy 301 TVGMAKLDPSSQALQALPYDPEMEEDSYDSFGSPSYPEVFPPLTGYGPEEEERGV 360
 Db 301 TVGMAKLDPSSQALQALPYDPEMEEDSYDSFGSPSYPEVFPPLTGYGPEEEERGV 360

Qy 361 KLGLGDFIFYSVLVGKAATGSGDNTTILACFVAILIGLCTLLLLAVFKKALPALPISI 420
 Db 361 KLGLGDFIFYSVLVGKAATGSGDNTTILACFVAILIGLCTLLLLAVFKKALPALPISI 420

Qy 421 TFGLIIFYSTDLNLRPFMDTLASHOLYI 448
 Db 421 TFGLIIFYSTDLNLRPFMDTLASHOLYI 448

RESULT 7
 W28508 standard; Protein: 447 AA.
 AC W28508;
 DT 07-DEC-1997 (first entry)
 DE Full AD/AD3LP sequence.
 KW AD3; AD4/AD3LP; Alzheimer's disease; chromosome; missegregation;
 KW presenilin; inhibitor; AD; trisomy 21; ss.
 OS Homo sapiens.
 PN W09707213-A2.
 PD 27-FEB-1997.
 PF 15-AUG-1996; U13314.
 PR 16-AUG-1995; US-002448.
 PA (HARD) HARVARD COLLEGE.
 PI Li J, Potter H;
 DR WPI: 97-165297/15.
 DR N-PSDB: T87426.
 PT Identifying genes which cause chromosome missegregation - useful for
 PT identifying causes of and treatments for diseases, e.g. Alzheimer's
 PT disease, cancer and ageing
 PS Claim 29; Fig 29; 77pp; English.
 CC Identifying genes which cause improper chromosome segregation,
 CC screening for inhibitors of chromosome missegregation and processes

CC caused by genes encoding chromosome missegregation promoters
 CC was exemplified using Alzheimer's disease. The sequences
 CC given in T87401 to T87426 can be used in the above methods.
 CC It is not clear from the figure legend, the figure and the
 CC disclosure of the specification which sequence of Fig 1 and Fig 28
 CC is the AD4/AD3LP or the AD3 sequence.
 SQ Sequence 447 AA;

Query Match 99.3%; Score 2320.5; DB 1; Length 447;
 Best Local Similarity 99.8%; Pred. No. 3.1e-235;
 Matches 447; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy 1 MLTFMADSEEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQRWSENEDEEDP 60
 Db 1 MLTFMADSEEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQRWSENEDEEDP 60

Qy 61 DRYVCSGVPGRPPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120
 Db 61 DRYVCSGVPGRPPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120

Qy 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLL 180
 Db 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLL 180

Qy 181 FLFTYIYLGELVKTYNVAMDYPTLLTWNFGAVGMVCIHWKGPLVLOQAYLIMISALMA 240
 Db 181 FLFTYIYLGELVKTYNVAMDYPTLLTWNFGAVGMVCIHWKGPLVLOQAYLIMISALMA 240

Qy 241 LVFTIKYLPENSAWVILGAISSYDVLVAVLCPKGPLRMLVETAQERNEPIFPALYSSAMVW 300
 Db 241 LVFTIKYLPENSAWVILGAISSYDVLVAVLCPKGPLRMLVETAQERNEPIFPALYSSAMVW 300

Qy 301 TVGMAKLDPSSQALQALPYDPEMEEDSYDSFGSPSYPEVFPPLTGYGPEEEERGV 360
 Db 301 TVGMAKLDPSSQALQALPYDPEMEEDSYDSFGSPSYPEVFPPLTGYGPEEEERGV 360

Qy 361 KLGLGDFIFYSVLVGKAATGSGDNTTILACFVAILIGLCTLLLLAVFKKALPALPISI 420
 Db 361 KLGLGDFIFYSVLVGKAATGSGDNTTILACFVAILIGLCTLLLLAVFKKALPALPISI 420

Qy 421 TFGLIIFYSTDLNLRPFMDTLASHOLYI 448
 Db 420 TFGLIIFYSTDLNLRPFMDTLASHOLYI 447

RESULT 8
 W05766 standard; Protein: 414 AA.
 AC W05766;
 DT 25-JUL-1997 (first entry)
 DE Presenilin-2 delta263-296 mutation.
 KW Presenilin-2; human; hps1-1; hps1-2; ps-2; integral membrane protein; AD;
 KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;
 KW depression; antibody; gene expression modulator; therapy; mutein.
 OS Homo sapiens.
 FH Key
 FT Location/Qualifiers
 FT misc_difference 263..264
 FT /note= "site of 34 residue deletion"
 PN W09634099-A2.
 PD 31-OCT-1996.
 PF 29-APR-1996; CA0263.
 PR 28-APR-1995; US-431048.
 PR 28-JUN-1995; US-496841.
 PR 31-JUL-1995; US-509359.
 PA (HSCR-) HSC RES & DEV LP.
 PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
 PI Fraser PE, Rommens JM, St George-Hyslop PH;
 DR WPI: 96-497631/49.
 PT New presenilin genes - useful for diagnosis, therapy and drug
 PT screening of familial Alzheimer's disease, cerebral disorders, etc.
 PS Claim 4; Page -; 178pp; English.
 CC W05763-W05766 represent mutated versions of the human presenilin-2

CC coding sequence for wild type human PS-2. The presenilins are a family of
CC highly conserved integral membrane proteins with a common structural
CC motif, common alternate splicing patterns, and common mutational hot spot
CC regions. Mutations in PS genes are implicated in familial Alzheimer's
CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
CC schizophrenia, depression etc., so detection of mutations in the DNA
CC encoding these sequences can be used for diagnosis of these diseases.
CC These proteins, or vectors that express them or containing antisense
CC sequences, antibodies selective for mutant forms of these proteins (such
CC as W05736) and modulators of PS gene expression are potentially useful
CC for treatment of AD etc. Transgenic animals are useful as models for drug
CC screening. The antibodies can also be used e.g. for affinity purification
CC and in immunoassays.
SQ Sequence 467 AA;

Query Match 62.8%; Score 1468; DB 1; Length 467;
Best Local Similarity 64.0%; Pred. No. 9.9e-146;
Matches 299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;
QY 24 ESPTPRSCQEGRGQDGCNTAQRWQSENEEDGEDDPDRYVCSGVP----GRPPG----- 75
Db 3 EIPAPLSTVFQNMSEDSSAIRSQNSQDSQERQOQHDRLNDPFIENGRPQSNRSQV 62
QY 75 -----LEELTKYGAHVIMLFPVPTLCMIVVYVATIKSVFYTEKNGOLIYPTPTEDT 128
Db 63 VEQDEEDELTKYGAHVIMLFPVPTLCMIVVYVATIKSVFYTEKNGOLIYPTPTEDT 122
QY 129 PSVQGRLLNSVLTLMISVIYVIMFLVVLVYKYCYKFIHGLWLSMLFLFYIYL 188
Db 123 ETVGORALHSILNAAIMISVIVIMTLLVVLVYKYCYKFIHGLWLSMLFLFYIYL 182
QY 189 GEVLKTYNVAMDYPLLTLTVNFGAVGVCIIHWKGLPLVLOQAYLIMISALMALVFKIYLP 248
Db 183 GEVFTKTYNVAMDYPLLTLTVNFGAVGVCIIHWKGLPLVLOQAYLIMISALMALVFKIYLP 242
QY 249 ENSAWVILGAISVDYDLVAVLCPKGLPMLVETAQERNEPFIYALYSSAMVTVGNKLD 308
Db 243 ENTAWLILAVISVDYDLVAVLCPKGLPMLVETAQERNEPFIYALYSSAMVTVGNKLD 302
QY 309 PSSQALQLPYDEME-----EDSYDSFGEPSPYVEFPEPLTGYPG----- 350
Db 303 PEAQ--RRVPKPKYNTQRAERETQDSGSDGSGFSEWEAQRDHGLPHRSTPESRAA 360
QY 350 -EEL-----EEERGVKLGIDPIFYSLVGRKAAATGSGDWTTLACFVAILGLCL 401
Db 361 VOELSGSILTSEDPEERGKVLGLGDFIFYSLVGRKAAATGSGDWTTLACFVAILGLCL 420
QY 402 TLLLAFAKALPALPISITFTGLIFVFTDNLVRFPMDFLASHQLYI 448
Db 421 TLLLAFAKALPALPISITFTGLIFVFTDNLVRFPMDFLASHQLYI 467

RESULT 11
W23966
ID W23966 standard; Protein; 467 AA.
AC W23966;
DE Mouse presenilin-1 homologue.
KW Presenilin-1; PS1 gene; mouse; familial Alzheimer's disease; FAD;
KW cerebral haemorrhage; schizophrenia; depression; epilepsy;
KW mental retardation; diagnosis; therapy; transgenic animal.
OS Mus musculus.
FH Key Location/Qualifiers
FT Domain 82..100
FT /label= TM1
FT /note= "transmembrane domain 1"
FT 101..132
FT /label= TM1-2
FT /note= "hydrophilic loop"
FT 133..154
FT /label= TM2
FT /note= "transmembrane domain 2"

FT Domain 155..163
FT /label= TM2-3
FT /note= "hydrophilic loop"
FT Domain 164..183
FT /label= TM3
FT /note= "transmembrane domain 3"
FT Domain 184..194
FT /label= TM4
FT /note= "hydrophilic loop"
FT Domain 195..212
FT /label= TM4
FT /note= "transmembrane domain 4"
FT Domain 213..220
FT /label= TM4-5
FT /note= "hydrophilic loop"
FT Domain 221..238
FT /label= TM5
FT /note= "transmembrane domain 5"
FT Domain 239..243
FT /label= TM5-6
FT /note= "hydrophilic loop"
FT Domain 244..262
FT /label= TM6
FT /note= "transmembrane domain 6"
FT Domain 263..407
FT /label= TM6-7
FT /note= "hydrophilic loop"
FT Domain 408..428
FT /label= TM8
FT /note= "transmembrane domain 8"
FT Misc_difference 177
FT /note= "Phe177Ser mutation site (Claim 1)"
FT Misc_difference 439
FT /note= "Ile439Val mutation site (Claim 1)"
FT W09801549-A2.
PD 15-JAN-1998.
PF 04-JUL-1997; CA0475.
PR 02-JAN-1997; US-034590.
PR 05-JUL-1996; US-021673.
PR 12-JUL-1996; US-021700.
PR 08-NOV-1996; US-029895.
PA (HSCR-) HSC RES & DEV LP
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
DR WPI; 98-286355/25.
PT New isolated mutant presenilin-1 genes - useful for developing
PT products for use in detection, diagnosis and therapy of Alzheimer's
PT disease and for drug screening
PS Disclosure; Page 199-200; 238pp; English.
CC This polypeptide comprises the murine presenilin-1 (PS1) homologue.
CC Its amino acid sequence was deduced from an isolated cDNA clone
CC (see W04668). Mutations in the human PS1 and PS2 genes (see
CC W04666-68) have been linked to the development in humans of forms
CC of familial Alzheimer's disease (FAD). All amino acids that are
CC mutated in analysed FAD pedigrees (see W23964) were conserved in
CC the murine homologue. Use of the nucleic acids and proteins
CC comprising or derived from presenilins can be made in screening and
CC diagnosing FAD, identifying and developing therapeutics for
CC treatment of FAD, and in producing cell lines and transgenic
CC animals useful as models of FAD. Methods for identifying
CC substances that bind to, or modulate the activity of a presenilin
CC protein, and methods for identifying substances that affect the
CC interaction of a presenilin-interacting protein with a presenilin
CC protein are also disclosed.
SQ Sequence 467 AA;

Query Match 62.8%; Score 1468; DB 1; Length 467;
Best Local Similarity 64.0%; Pred. No. 9.9e-146;
Matches 299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;
QY 24 ESPTPRSCQEGRGQDGCNTAQRWQSENEEDGEDDPDRYVCSGVP-----GRPPG----- 75

CC peptides. Antisense nucleic acid may also be used to regulate expression
CC of the P51/429 gene, and both nucleic acids and peptides are useful as
CC size markers in electrophoresis, chromatography etc. The transgenic
CC animals are used as models for AD, e.g. for testing drugs. Regulators of
CC the P51/429 gene or polypeptide can be used to treat e.g. AD or diseases
CC involving mitochondrial pathology, apoptosis and neurodegeneration.
CC Typical regulators are antisense sequences, ribozymes, aptamers,
CC synthetic or natural compounds. (ii) may also be used to target other
CC coding sequences to particular cellular locations.
SQ Sequence 467 AA;

Query Match 62.8%; Score 1467; DB 1; Length 467;
Best Local Similarity 65.7%; Pred. No. 1.3e-145;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;
QY 24 ESTPTSCGEGRGPEGENTAGWRQSENEDEEDDPDRVVCVGVP-----GRPPG--- 75
DQ 3 ELPAPLSYFQNAQMSDNHLSNTVRSONRQRHNDRR-SLGHPEPLSGRPPQGNRSRQ 61
QY 75 -----LEEELTKYGAHVIMLFVPTVLCMIVVATIKSVFYTEKNGQLIYTPPTED 127
DQ 62 VVEQDEEDELTKYGAHVIMLFVPTVLCMIVVATIKSVFSFYTRKDGQLIYTPPTED 121
QY 128 TPVGORLLNSVLTLMISIVIVMTIFLVLYKYRCYKTHGWLIMSSLMLLFLFYIY 187
DQ 122 TETVGORALHSILNAIMISIVIVMTIFLVLYKYRCYKVIHAWLISSLLLLFFFSFIY 181
QY 188 LGEVLTYNVADYPILLTLFWNFGVMYCIHWKGPLVQQAYLYMISALMALVFIKYL 247
DQ 182 LGEVFTYNVADYITVALLIWNFGVGMISHWKGPLRQQAYLYMISALMALVFIKYL 241
QY 248 PEWSANVILGAIISVDLVAVLCPLKPLMLVETAQERNPIFIPALIISSAMVTVGVMAKL 307
DQ 242 PEWTALILAVISVDLVAVLCPLKPLMLVETAQERNETLFPALIISSAMVTVGVMAEG 301
QY 308 DPSSQAL--QLPYDPE-MEEDSVDSFGE--PSYPVEEPPLTGYPG----- 350
DQ 302 DPAQRVSKNSKYNKSTRESQDITVAENDDGGFSEWAEQADSHUGPHRSTPESRAAV 361
QY 350 EEL-----EEERGVKGLGDFIFYSVLVGAATGSGDWNNTLACFVAILIGLICLT 402
DQ 362 QELSSSILAGEDPEERGVKGLGDFIFYSVLVGAATGSGDWNNTLACFVAILIGLICLT 421
QY 403 LLLAVFKKALPALPISITFLGYFSTDNLPVFPMDTTLASHQLIY 448
DQ 422 LLLAIFKKALPALPISITFLGYFATDYLVQPFMDQLAFHOFYI 467
RESULT 14
W23964
ID W23964 standard; Protein; 467 AA.
AC W23964;
DT 20-JUL-1998 (first entry)
DE Human presenilin-1.
KW Presenilin-1; P51 gene; human; familial Alzheimer's disease; FAD;
KW cerebral haemorrhage; schizophrenia; depression; epilepsy;
KW mental retardation; diagnosis; therapy; transgenic animal.
OS Homo sapiens.
FH Key
FT Location/Qualifiers
FT 82..100
FT /label= TM1
FT /note= "transmembrane domain 1"
FT 101..132
FT /label= TM1-2
FT /note= "hydrophilic loop"
FT 133..154
FT /label= TM2
FT /note= "transmembrane domain 2"
FT 155..163
FT /label= TM2-3
FT /note= "hydrophilic loop"
FT 164..183

FT /label= TM3
FT /note= "transmembrane domain 3"
FT 184..194
FT /label= TM3-4
FT /note= "hydrophilic loop"
FT 195..212
FT /label= TM4
FT /note= "transmembrane domain 4"
FT 213..220
FT /label= TM4-5
FT /note= "hydrophilic loop"
FT 221..238
FT /label= TM5
FT /note= "transmembrane domain 5"
FT 239..243
FT /label= TM5-6
FT /note= "hydrophilic loop"
FT 244..262
FT /label= TM6
FT /note= "transmembrane domain 6"
FT 263..407
FT /label= TM6-7
FT /note= "hydrophilic loop"
FT 408..428
FT /label= TM8
FT /note= "transmembrane domain 8"
FT Misc_difference 177 /note= "Phel17Ser mutation site (Claim 1)"
FT Misc_difference 439 /note= "Ile439Val mutation site (Claim 1)"
FT Misc_difference 257 /note= "Asp257Ala mutation site, associated with
FT residue 258-290 deletion (Claim 1)"
FT Misc_difference 258..290 /note= "residue 258-290 deletion mutant, associated
FT with Asp257Ala mutation (Claim 1)"
FT Misc_difference 143 /note= "Ile143Thr mutation site (Claim 18)"
FT Misc_difference 146 /note= "Met146Leu mutation site (Claim 18)"
FT Misc_difference 171 /note= "Leu171Pro mutation site (Claim 18)"
FT Misc_difference 260 /note= "Ala260Val mutation site (Claim 18)"
FT Misc_difference 263 /note= "Cys263Arg mutation site (Claim 18)"
FT Misc_difference 264 /note= "Pro264Leu mutation site (Claim 18)"
FT Misc_difference 267 /note= "Pro267Ser mutation site (Claim 18)"
FT Misc_difference 280 /note= "Glu280Ala mutation site (Claim 18)"
FT Misc_difference 280 /note= "Glu280Gly mutation site (Claim 18)"
FT Misc_difference 285 /note= "Ala285Val mutation site (Claim 18)"
FT Misc_difference 286 /note= "Leu286Val mutation site (Claim 18)"
FT Misc_difference 322 /note= "Leu322Val mutation site (Claim 18)"
FT Misc_difference 392 /note= "Leu392Val mutation site (Claim 18)"
FT Misc_difference 410 /note= "Cys410Tyr mutation site (Claim 18)"
FT Misc_difference 79 /note= "Ala79Xaa mutation site"
FT Misc_difference 82 /note= "Val82Leu mutation site"
FT Misc_difference 96 /note= "Val96Phe mutation site"
FT Misc_difference 115 /note= "Tyr115His mutation site"
FT Misc_difference 139

Matches 305; Conservative 40; Mismatches 79; Indels 42; Gaps 8;

```
QY 24 ESTPTSCQGGQGPDEGNTAQRSGEENEDGEDPDYVCSGVP-----GRPPG---- 75
Db 3 ELAPLSYFQNAQMSDNHLSNVRSDNRRERQEHDRR-SLGHPEPLSNGRPQGNRQ 61
QY 75 -----LEELTLKYGAKHVIMLFVPTLCMIVVATIKSVREYTKNGQLIYPTFTED 127
Db 62 VEQDEDEDELTLKYGAKHVIMLFVPTLCMIVVATIKSVFYTRKDGOLIYPTFTED 121
QY 128 TPSVGQRLNSVNTLIMISIVVMTFLVLYKYRCYKFIHGWLMSSMLLFLFYIY 187
Db 122 TETVGQALHSILNAAIMISIVVMTLLVLYKYRCYKVIHAWLIISLLEFFSFIY 181
QY 188 LGEVLKTYNYAMDYPTLLLTWNFGAVGWCIIHWKGPLVLQQAAYLIMISALMALVFIKYL 247
Db 182 LGEVFKTYNAVDTITVALLIWNFGVVGMSIHWKGPLRLQQAAYLIMISALMALVFIKYL 241
QY 248 PEWSAWILGAISYIDLAVLCPKPLRMLVETAQERNEIPFPALIIYSSAMVTVGMAKL 307
Db 242 PENTAWILLAVISYIDLAVLCPKPLRMLVETAQERNEILFPAVIYSSMVLVYNAEG 301
QY 308 DPSSQAL--QLPYDPE-MEEDSYDSFGE---PSYPEVFEPPLTGYPG----- 350
Db 302 DPEAQRVSKNSKYNASTERESQDTVAENDDGGFSEWEAQRDSHLGPHRSTPESRAAV 361
QY 350 EEL-----EEEEGVKLGDFIYSVLVKAATGSGDWNTTACFVAILIGLCLT 402
Db 362 QELSSSILAGEDPEERGKVLGLDFIYSVLVKAATGSGDWNTTACFVAILIGLCLT 421
QY 403 LLLIAVFKKALPALPISITFGLIFYFSTDNLVRPFMDTLASHOLYI 448
Db 422 LLLIAVFKKALPALPISITFGLVYFATDYLVPFMDQLAFHQFYI 467
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Search completed: March 20, 2000, 05:31:20
Job time: 4209 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 19:01:42 ; Search time 26.47 Seconds
(without alignments)
225.059 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFMADSEEVCDERTSL.....STDNLVRPFMDTIAHQLYI 448

Scoring table: BLOSUM62

Searched: 133990 seqs, 13297546 residues

Database : Issued_Patents_AA.*

Word size : 0

Number of hits that pass the threshold : 133990
1: /cgn2.6/ptodata/2/iaa/5A_COMB.pep.*
2: /cgn2.6/ptodata/2/iaa/5B_COMB.pep.*
3: /cgn2.6/ptodata/2/iaa/6_COMB.pep.*
4: /cgn2.6/ptodata/2/iaa/PCTUS9_COMB.pep.*
5: /cgn2.6/ptodata/2/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	2336	100.0	448	2	US-08-967-101-137
2	2336	100.0	448	2	US-08-592-541-137
3	2320.5	99.3	447	2	US-08-875-972-29
4	1923	82.3	372	2	US-08-967-101-138
5	1923	82.3	372	2	US-08-592-541-138
6	1623.5	69.5	376	2	US-08-875-972-2
7	1457	62.8	467	2	US-08-967-101-134
8	1467	62.8	467	2	US-08-592-541-134
9	1461	62.5	467	3	US-08-670-964-2
10	1461	62.5	467	2	US-08-967-101-2
11	1461	62.5	467	2	US-08-592-541-2
12	1434	62.2	463	3	US-08-670-964-4
13	1438	61.6	407	2	US-08-875-972-4
14	1437	61.5	463	2	US-08-670-479-18
15	1418	60.7	467	2	US-08-367-101-4
16	1418	60.7	467	2	US-08-592-541-4
17	1150	49.2	541	2	US-08-967-101-166
18	1150	49.2	541	2	US-08-592-541-166
19	110	4.7	1294	2	US-08-819-288-3
20	108	4.6	1321	1	US-08-261-822A-3
21	108	4.6	1321	4	PCT-US95-07744A-3
22	101	4.3	1334	2	US-08-986-545-2
23	100	4.3	400	1	US-08-602-010A-8
24	100	4.3	400	1	US-08-680-726A-8
25	95	4.1	391	1	US-07-816-283-2
26	95	4.1	391	1	US-08-417-103-4
27	93	4.0	391	1	US-07-816-283-2
28	93	4.0	391	1	US-08-417-103-2
29	93	4.0	391	1	US-08-417-103-14
30	91	3.9	509	2	US-09-031-392-6
31	88.5	3.8	452	1	US-08-117-361C-1
32	88.5	3.8	3169	2	US-08-477-451-6
33	87	3.7	492	2	US-08-355-844-3
34	87	3.7	492	4	PCT-US95-16126-3

ALIGNMENTS

RESULT 1

US-08-967-101-137

: Sequence 137, Application US/08967101

: Patent No. 5840540

: GENERAL INFORMATION:

: APPLICANT: ST. GEORGE-HYSLOP, PETER H

: APPLICANT: ROMMENS, JOHANNA M

: APPLICANT: FRASER, PAUL E

: TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED

: TITLE OF INVENTION: TO ALZHEIMER'S DISEASE

: NUMBER OF SEQUENCES: 183

: CORRESPONDENCE ADDRESS:

: ADDRESSEE: TESTA, HURWITZ & THIBEAULT

: STREET: High Street Tower - 125 High Street

: CITY: Boston

: STATE: Massachusetts

: COUNTRY: U.S.A.

: ZIP: 02110

: COMPUTER READABLE FORM:

: MEDIUM TYPE: Floppy disk

: COMPUTER: IBM PC compatible

: OPERATING SYSTEM: PC-DOS/MS-DOS

: SOFTWARE: Patentin Release #1.0, Version #1.30

: CURRENT APPLICATION DATA:

: APPLICATION NUMBER: US/08/967,101

: FILING DATE: 10-NOV-1997

: CLASSIFICATION: 435

: PRIOR APPLICATION DATA:

: APPLICATION NUMBER: 08/592,541

: FILING DATE:

: ATTORNEY/AGENT INFORMATION:

: NAME: Pitcher, Edmund R.

: TELECOMMUNICATION INFORMATION:

: TELEPHONE: (617) 248-7000

: TELEFAX: (617) 248-7100

: INFORMATION FOR SEQ ID NO: 137:

: SEQUENCE CHARACTERISTICS:

: LENGTH: 448 amino acids

: TYPE: amino acid

: STRANDEDNESS: single

: TOPOLOGY: linear

: MOLECULE TYPE: protein

US-08-967-101-137

Query Match 100.0%; Score 2336; DB 2; Length 448;

Best Local Similarity 100.0%; Pred. No. 1e-231;

Matches 448; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MLTFMADSEEVCDERTSLMSAESPTPRSCQGRQGPEDGNTAQWRSQNEEDGEDP 60

DB 1 MLTFMADSEEVCDERTSLMSAESPTPRSCQGRQGPEDGNTAQWRSQNEEDGEDP 60

QY 61 DRYVCSGVPGPPGLEELTLKYGAKHVMFLPVPVTLGMVVVATIKSVRFYTERKNGQLI 120

DB 61 DRYVCSGVPGPPGLEELTLKYGAKHVMFLPVPVTLGMVVVATIKSVRFYTERKNGQLI 120

US-08-875-972-29

Query Match 99.3%; Score 2320.5; DB 2; Length 447;
Best Local Similarity 99.8%; Pred. No. 4e-230;
Matches 447; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 MLTMSDSEEEVCDERTSLMSASPTPRSCQGRQGPEDGENTAQRNEEDGEDP 60
DB 1 MLTMSDSEEEVCDERTSLMSASPTPRSCQGRQGPEDGENTAQRNEEDGEDP 60
QY 61 DRYVCSGVPGRPGLEELIKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120
DB 61 DRYVCSGVPGRPGLEELIKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120
QY 121 YTPETDTPSGVORLNSVNTLMISIVVMTIFLVLYKYRCYKFTGWLIMSSMLL 180
DB 121 YTPETDTPSGVORLNSVNTLMISIVVMTIFLVLYKYRCYKFTGWLIMSSMLL 180
QY 181 FLFTYIYLGVLKTYNVAMDPTLLTWNFGAVGVCIHMKGPLVLQOAYLIMISALMA 240
DB 181 FLFTYIYLGVLKTYNVAMDPTLLTWNFGAVGVCIHMKGPLVLQOAYLIMISALMA 240
QY 241 LVFTIKYLPWSAWVILGAISYIDLAVLCPKGPLRMLVETAQRNEPIFPALYSSAMVW 300
DB 241 LVFTIKYLPWSAWVILGAISYIDLAVLCPKGPLRMLVETAQRNEPIFPALYSSAMVW 300
QY 301 TVGMAKLDPSOGALQLPYPDEEDSDYSGEPEYFEPPLTGYGPELEEEERGV 360
DB 301 TVGMAKLDPSOGALQLPYPDEM-EDSYDSGEPEYFEPPLTGYGPELEEEERGV 359
QY 361 KLGIGDFIFYSVLVGRKAAATGSGDWNNTLACFVAILGICLTLTLLAVFKKALPALPISI 420
DB 360 KLGIGDFIFYSVLVGRKAAATGSGDWNNTLACFVAILGICLTLTLLAVFKKALPALPISI 419
QY 421 TFGLIYFYSTDLNVRPMDTLASHQLYI 448
DB 420 TFGLIYFYSTDLNVRPMDTLASHQLYI 447

RESULT 4

US-08-967-101-138
Sequence 138, Application US/08967101
Patent No. 5840540

GENERAL INFORMATION:

APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:

ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,101
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/592,541

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Pitcher, Edmund R.

TELECOMMUNICATION INFORMATION:

TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 138:
SEQUENCE CHARACTERISTICS:
LENGTH: 372 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-967-101-138

Query Match 82.3%; Score 1923; DB 2; Length 372;
Best Local Similarity 100.0%; Pred. No. 2e-189;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 77 BELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLIYTPFTEDTPSVQORLL 136
DB 1 BELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLIYTPFTEDTPSVQORLL 60
QY 137 NSVLTLMISIVVMTIFLVLYKYRCYKFTGWLIMSSMLLFTYIYLGVLKTYN 196
DB 61 NSVLTLMISIVVMTIFLVLYKYRCYKFTGWLIMSSMLLFTYIYLGVLKTYN 120
QY 197 VAMDPTLLTWNFGAVGVCIHMKGPLVLQOAYLIMISALMALVFIKYLPWSAWVIL 256
DB 121 VAMDPTLLTWNFGAVGVCIHMKGPLVLQOAYLIMISALMALVFIKYLPWSAWVIL 180
QY 257 GAISYIDLAVLCPKGPLRMLVETAQRNEPIFPALYSSAMVWTVGMAKLDPSOGALQ 316
DB 181 GAISYIDLAVLCPKGPLRMLVETAQRNEPIFPALYSSAMVWTVGMAKLDPSOGALQ 240
QY 317 LPYDPEMEDSDYSGEPEYFEPPLTGYGPELEEEERGVKLGDFIFYSVLVGR 376
DB 241 LPYDPEMEDSDYSGEPEYFEPPLTGYGPELEEEERGVKLGDFIFYSVLVGR 300
QY 377 AAATGSGDWNNTLACFVAILGICLTLTLLAVFKKALPALPISITFGLIYFYSTDLNVRP 436
DB 301 AAATGSGDWNNTLACFVAILGICLTLTLLAVFKKALPALPISITFGLIYFYSTDLNVRP 360
QY 437 FMDTLASHQLYI 448
DB 361 FMDTLASHQLYI 372

RESULT 5

US-08-592-541-138
Sequence 138, Application US/08592541
Patent No. 5986054

GENERAL INFORMATION:

APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:

ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/592,541
FILING DATE:

CLASSIFICATION: 800

ATTORNEY/AGENT INFORMATION:

SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,101
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 134:
SEQUENCE CHARACTERISTICS:
LENGTH: 467 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-967-101-134

Query Match 62.8%; Score 1467; DB 2; Length 467;
Best Local Similarity 65.7%; Pred. No. 2e-142;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;

QY 24 ESPTRSCQGRGQPEDGNTAQWRSQNEDEEDDPDRYVCSGVP-----GRPPG---- 75
DB 3 ELPAISYFQNAQMSDNHLSNTRVSDNDRERQEHDRR-SLGHPEPLSNRQGNRSRQ 61
QY 75 -----LEELILKYGAKHVIMLFVPTLCMVVVVATIKSVFYTKNGQLIYPTFTED 127
DB 62 VVEQDEEDELTKYGAHVIMLFVPTLCMVVVVATIKSVFYTKNGQLIYPTFTED 121
QY 128 TPSVGORLNSVLNTLMISVIVVMTIFLVLYKYRCYKFIHGWLMSSLMILFLFTYIY 187
DB 122 TETVGORALHSILNAIMISVIVVMTIFLVLYKYRCYKFIHAWLISSLLLFYFSFIY 181
QY 188 LGEVLTNYVNDYPTLLTVNFGAVGVCIHKGPLVQLQAYLIMISALMALVFIKYL 247
DB 182 LGEVFTYNAVVDYITVALLIWNFGVGMISIHKGPLRLQQAAYLIMISALMALVFIKYL 241
QY 248 PEWSAVILGALSYVDLVAVLCPKPLRMVETAQRNEPIFPALIIYSSAMVTVGMAKL 307
DB 242 PEWTAWLILAVISVDLVAVLCPKPLRMVETAQRNETLFPALIIYSSAMVTVGMAL 301
QY 308 DPSSQAGL--QLPYDPE-MEEDSYDSFGE---PSYPEVPEPLTGYPG----- 350
DB 302 DPEAQRVSKNSKYNASTERSQDTVAENDDGGFSEWEAQRDHLGPHRSTPESRAAV 361
QY 350 EEL-----EEEEERGVKLGDFIFYSVLVGAATGSDWNTTACFVAILIGLCLT 402
DB 362 QELSSSILAGEDPEERGVKLGDFIFYSVLVGAATGSDWNTTACFVAILIGLCLT 421
QY 403 LLLAVFKKALPALPISITFTGLIFYSTDLNLRPFMDTLASHQLYI 448
DB 422 LLLAIFKKALPALPISITFTGLVYFATDYLQVPMQDLAFHOFYI 467

RESULT 8
US-08-592-541-134
Sequence 134, Application US/08592541
Patent No. 5986054
GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183
CORRESPONDENCE ADDRESS:
ADDRESSEE: TESTA, HURWITZ & THIBEAULT

STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/592,541
FILING DATE:
CLASSIFICATION: 800
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 134:
SEQUENCE CHARACTERISTICS:
LENGTH: 467 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-592-541-134

Query Match 62.8%; Score 1467; DB 2; Length 467;
Best Local Similarity 65.7%; Pred. No. 2e-142;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;

QY 24 ESPTRSCQGRGQPEDGNTAQWRSQNEDEEDDPDRYVCSGVP-----GRPPG---- 75
DB 3 ELPAISYFQNAQMSDNHLSNTRVSDNDRERQEHDRR-SLGHPEPLSNRQGNRSRQ 61
QY 75 -----LEELILKYGAKHVIMLFVPTLCMVVVVATIKSVFYTKNGQLIYPTFTED 127
DB 62 VVEQDEEDELTKYGAHVIMLFVPTLCMVVVVATIKSVFYTKNGQLIYPTFTED 121
QY 128 TPSVGORLNSVLNTLMISVIVVMTIFLVLYKYRCYKFIHGWLMSSLMILFLFTYIY 187
DB 122 TETVGORALHSILNAIMISVIVVMTIFLVLYKYRCYKFIHAWLISSLLLFYFSFIY 181
QY 188 LGEVLTNYVNDYPTLLTVNFGAVGVCIHKGPLVQLQAYLIMISALMALVFIKYL 247
DB 182 LGEVFTYNAVVDYITVALLIWNFGVGMISIHKGPLRLQQAAYLIMISALMALVFIKYL 241
QY 248 PEWSAVILGALSYVDLVAVLCPKPLRMVETAQRNEPIFPALIIYSSAMVTVGMAKL 307
DB 242 PEWTAWLILAVISVDLVAVLCPKPLRMVETAQRNETLFPALIIYSSAMVTVGMAL 301
QY 308 DPSSQAGL--QLPYDPE-MEEDSYDSFGE---PSYPEVPEPLTGYPG----- 350
DB 302 DPEAQRVSKNSKYNASTERSQDTVAENDDGGFSEWEAQRDHLGPHRSTPESRAAV 361
QY 350 EEL-----EEEEERGVKLGDFIFYSVLVGAATGSDWNTTACFVAILIGLCLT 402
DB 362 QELSSSILAGEDPEERGVKLGDFIFYSVLVGAATGSDWNTTACFVAILIGLCLT 421
QY 403 LLLAVFKKALPALPISITFTGLIFYSTDLNLRPFMDTLASHQLYI 448
DB 422 LLLAIFKKALPALPISITFTGLVYFATDYLQVPMQDLAFHOFYI 467

RESULT 9
US-08-670-964-2
Sequence 2, Application US/08670964
Patent No. 6010874
GENERAL INFORMATION:
APPLICANT: Hardy, John A.
TITLE OF INVENTION: EARLY ONSET ALZHEIMER'S DISEASE

QY 308 DPSSQAL--QLPYDPE-MEEDSYDSFGE---PSYPEVEFPEPLTGYPG----- 350
 Db 302 DPEARRVSKNSKYNASTERESQDTVAENDGGFSEWEAQRSHLGPHERSTPESRAV 361
 QY 350 EEL-----EEERGVKGLGDFIFYSVLVGRKAATGSGDWNNTLACFVAILIGLCLT 402
 Db 362 QELSSSILAGEDPEERGKVLGDFIFYSVLVGRASATASGDWNNTIACFVAILIGLCLT 421
 QY 403 LLLAVFKKALPALPISITFGLIFFSTDNLVPRFMDTLASHQLYI 448
 Db 422 LLLAIFKKALPALPISITFGLVFYFATDYLVPQFMDQLAFHQFYI 467

RESULT 11
 US-08-592-541-2
 ; Sequence 2, Application US/08592541
 ; Patent No. 5986054
 ; GENERAL INFORMATION:
 ; APPLICANT: ST. GEORGE-HYSLOP, PETER H
 ; APPLICANT: ROMMENS, JOHANNA M
 ; APPLICANT: FRASER, PAUL E
 ; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 ; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
 ; NUMBER OF SEQUENCES: 183
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
 ; STREET: High Street Tower - 125 High Street
 ; CITY: Boston
 ; STATE: Massachusetts
 ; COUNTRY: U.S.A.
 ; ZIP: 02110
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/592,541
 ; FILING DATE:
 ; CLASSIFICATION: 800
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Pitcher, Edmund R.
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (617) 248-7000
 ; TELEFAX: (617) 248-7100
 ; INFORMATION FOR SEQ ID NO: 2:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 467 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ; US-08-592-541-2

Query Match 62.5%; Score 1461; DB 2; Length 467;
 Best Local Similarity 65.5%; Pred. No. 8.1e-142;
 Matches 305; Conservative 39; Mismatches 80; Indels 42; Gaps 8;

QY 24 ESPTPRSCQEGRQGPEDGENTAQWRSQNEEDGEDPDRYVCSGVP-----GRPPG----- 75
 Db 3 ELPAFLSYFQNAQSEDNHLSNTVRSQNDNREOEHDRR-SLGHPEPLSNGRPGQNSGR 61
 QY 75 -----LEELTLKYGAKHVIMLFVPVTLCLMIVVATIKSVFTEKNGOLITPFTED 127
 Db 62 VVEQDEEDELTLKYGAKHVIMLFVPVTLCLMIVVATIKSVFTRKDGOLITPFTED 121

QY 128 TPSYQORLLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGLMISLMLFFLTYYI 187
 Db 122 TETVQORALHSILNAIMISIVVMTILLVLYKYRCYKVIHAWLIISLMLFFLTYYI 181

QY 188 LGEVLKTYNVAMDPTLLLTWNFGAVGWCIVHWKGPLVQQAYLIMISALMALVFIKYL 247

Db 162 LCEVEKTYNVADYITVALLIWNLGIVGVGMISTHWKGPLRQQAYLIMISALMALVFIKYL 241
 QY 248 PEWSAWVILGAISVYDVLAVVLCPKGPLRMLVETAQERNPIFPALIISSAMVWTVGMAKL 307
 Db 242 PEWTAWLLAVISVYDVLAVVLCPKGPLRMLVETAQERNETLFPALIISSMTMVLVNAEG 301
 QY 308 DPSSQAL--QLPYDPE-MEEDSYDSFGE---PSYPEVEFPEPLTGYPG----- 350
 Db 302 DPEARRVSKNSKYNASTERESQDTVAENDGGFSEWEAQRSHLGPHERSTPESRAV 361
 QY 350 EEL-----EEERGVKGLGDFIFYSVLVGRKAATGSGDWNNTLACFVAILIGLCLT 402
 Db 362 QELSSSILAGEDPEERGKVLGDFIFYSVLVGRASATASGDWNNTIACFVAILIGLCLT 421
 QY 403 LLLAVFKKALPALPISITFGLIFFSTDNLVPRFMDTLASHQLYI 448
 Db 422 LLLAIFKKALPALPISITFGLVFYFATDYLVPQFMDQLAFHQFYI 467

RESULT 12
 US-08-670-964-4
 ; Sequence 4, Application US/08670964
 ; Patent No. 6010874
 ; GENERAL INFORMATION:
 ; APPLICANT: Hardy, John A.
 ; TITLE OF INVENTION: EARLY ONSET ALZHEIMER'S DISEASE
 ; TITLE OF INVENTION: GENE AND GENE PRODUCTS
 ; NUMBER OF SEQUENCES: 4
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: SmithKline Beecham Corporation
 ; STREET: 709 Swedeland Road - UW2220; P.O. Box 15
 ; CITY: Philadelphia
 ; STATE: PA
 ; COUNTRY: USA
 ; ZIP: 19406
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Diskette
 ; COMPUTER: IBM Compatible
 ; OPERATING SYSTEM: DOS
 ; SOFTWARE: FastSeq for Windows Version 2.0
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/670,964
 ; FILING DATE: 26-JUN-1996
 ; CLASSIFICATION: 435
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 60/001,142
 ; FILING DATE: 13-JUL-1995
 ; APPLICATION NUMBER: 60/001,501
 ; FILING DATE: 18-JUL-1995
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Han, William T
 ; REGISTRATION NUMBER: 34,344
 ; REFERENCE/DOCKET NUMBER: P50358
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: 610-270-5219
 ; TELEFAX: 610-270-5090
 ; TELEX:
 ; INFORMATION FOR SEQ ID NO: 4:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 463 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ; US-08-670-964-4

Query Match 62.2%; Score 1454; DB 3; Length 463;
 Best Local Similarity 65.3%; Pred. No. 4.2e-141;
 Matches 303; Conservative 40; Mismatches 79; Indels 42; Gaps 8;

QY 24 ESPTPRSCQEGRQGPEDGENTAQWRSQNEEDGEDPDRYVCSGVP---GRPPG----- 75

REFERENCE/DOCKET NUMBER: P50361
TELEPHONE: 610-270-5219
TELEFAX: 610-270-5030
TELEX:

INFORMATION FOR SEQ ID NO: 18:

SEQUENCE CHARACTERISTICS:
LENGTH: 463 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
HYPOTHETICAL: NO
ANTI-SENSE: NO
FRAGMENT TYPE: N-terminal
ORIGINAL SOURCE:
US-08-670-479-18

Query Match 61.5%; Score 1437; DB 2; Length 463;
Best Local Similarity 64.9%; Pred. No. 2.3e-139;
Matches 301; Conservative 41; Mismatches 80; Indels 42; Gaps 8;

QY 24 ESPTPRSCQGRQGEDGENTAQWRSQNEEDGEDDPDRYVCSGVP---GRPPG----- 75
DB 3 ELPAPLSYFQNAQMSDNHLS---NTNDRERQEHNDRRSLGHPEPLSNRGPQGNRSQV 59
QY 75 -----LEELILKYGAHVIMLVFVPTLCMVVATIKSVFYTEKNGOLIYTPPTEDTP 129
DB 60 EQDEEDELTKYGAHVIMLVFVPTLCMVVATIKSVFYTRKDGOLIYTPPTEDTE 119
QY 130 SVGORLLNSVLTLMISVIVMTILVLYKYRCYKIHWGLIMSSLMFLFYIYL 189
DB 120 TVGQALHSILNAALMISVIVMTILVLYKYRCYKIHWGLIISLLLFEEFYILG 179
QY 190 EYLKTYNVAMDYPTLLLVFNWFGAVGMYCIHWKGLVQLQAYLIMISALMALVFIKYLPE 249
DB 180 EVKTYNVAMDYPTLLLVFNWFGVGMISLHWKGLRQLQAYLIMISALMALVFIKYLPE 239
QY 250 WSAWVILGAI SYDLVAVLCPKGPLRMLVETAQERNPIFPALIIYSSAMVTVGMAKLDP 309
DB 240 WTAWLILAVISYDLVAVLCPKGPLRMLVETAQERDETLFPALIIYSSMTVMVLVNAEGDP 299
QY 310 SSQGLA--QLPYDPE--MEEDSYDFGE--PSYPVFEPPLTGYPG-----EE 351
DB 300 EAQRVSKNSYNAESTRESQDTVAENDDGGFSEWEAQRDHSLGPHRSTPESRAAVQE 359
QY 352 L-----EEEEERGVKLGIDGFIFYSVLVGKAAATGSDWNTTLACFVAILIGLCLTLL 404
DB 360 LSSSILAGEDPEERGVKLGIDGFIFYSVLVGKASATASGDWNTTLACFVAILIGLCLTLL 419
QY 405 LLAVFKKALPALPISITFGLIFYFSTDNLVRFPMDTLASHQLYI 448
DB 420 LLAIFKKALPALPISITFGLIFYFATDYLVPQFMDQLAFHQFYI 463

RESULT 15
US-08-967-101-4
Sequence 4, Application US/08967101
Patent No. 5840540
GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183
CORRESPONDENCE ADDRESS:
ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.

ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,101
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541

FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541

FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 467 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-967-101-4

Query Match 60.7%; Score 1418; DB 2; Length 467;
Best Local Similarity 61.9%; Pred. No. 2.1e-137;
Matches 289; Conservative 48; Mismatches 86; Indels 44; Gaps 6;

QY 24 ESPTPRSCQGRQGEDGENTAQWRSQNEEDGEDDPDRYVCSGVP---GRPPG----- 75
DB 3 ELPAPLSYFQNAQMSDNHLS---NTNDRERQEHNDRRSLGHPEPLSNRGPQGNRSQV 62
QY 75 -----LEELILKYGAHVIMLVFVPTLCMVVATIKSVFYTEKNGOLIYTPPTEDT 128
DB 63 EQDEEDELTKYGAHVIMLVFVPTLCMVVATIKSVFYTRKDGOLIYTPPTEDT 122
QY 129 PSVGORLLNSVLTLMISVIVMTILVLYKYRCYKIHWGLIMSSLMFLFYIYL 188
DB 123 EVGQALHSILNAALMISVIVMTILVLYKYRCYKIHWGLIISLLLFEEFYIYL 182
QY 189 GEVLKTYNVAMDYPTLLLVFNWFGAVGMYCIHWKGLVQLQAYLIMISALMALVFIKYL 248
DB 183 GEVFTYNVAMDYPTLLLVFNWFGVGMIAHWKGLRQLQAYLIMISALMALVFIKYL 242
QY 249 EWSAWVILGAI SYDLVAVLCPKGPLRMLVETAQERNPIFPALIIYSSAMVTVGMAKLD 308
DB 243 EWTAWLILAVISYDLVAVLCPKGPLRMLVETAQERNETLFPALIIYSSMTVMVLVNAEGD 302
QY 309 PSSQGLALQPYDPEME-----EDSYDFGEPSPYVFEPPPLTGYPG----- 350
DB 303 PEAQ--RRVPKPKYNTQRAERETQDSGSDNDGGFSEWEAQRDHSLGPHRSTPESRAA 360
QY 350 -EEL-----EEEEERGVKLGIDGFIFYSVLVGKAAATGSDWNTTLACFVAILIGLCL 401
DB 361 VQELSGSILTSDEPERGVKLGIDGFIFYSVLVGKASATASGDWNTTLACXVAILIGLCL 420
QY 402 TLLLLAVFKKALPALPISITFGLIFYFSTDNLVRFPMDTLASHQLYI 448
DB 421 XLLLLAIYKKGXPAPXISITFTGFGVFXFATDYLVPQFMDQLAFHQFYI 467

Search completed: March 18, 2000, 19:55:31
Job time: 3229 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 14:03:43 ; Search time 41.25 Seconds
(without alignments)
512.287 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFWASDSEEEVCDERTSL.....STDNLVRPFMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 142080 seqs, 47169319 residues

Database : PIR_62:*

Word size : 0
Number of hits that pass the threshold : 142080
1: pir1: *
2: pir2: *
3: pir3: *
4: pir4: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	2336	100.0	448	2 I58098	ES-1 protein - hum
2	2328	99.7	448	2 A56993	presenilin 2 - hum
3	2215.5	94.8	442	2 I39174	seven trans-membra
4	1624.5	69.5	449	2 JC5391	presenilin-beta -
5	1468	62.8	467	2 I78388	S182 protein - mou
6	1467	62.8	467	2 S58396	presenilin 1, spli
7	1454	62.2	463	2 S63883	presenilin I-463 -
8	1449.5	62.1	433	2 JC5390	presenilin-alpha -
9	1447	61.9	467	2 JC5080	presenilin 1 prote
10	1440	61.6	463	2 JC5081	presenilin 1 prote
11	1035.5	44.3	374	2 S63884	presenilin 1, spli
12	967	41.4	461	2 S60253	sel-12 protein - C
13	524.5	22.5	358	2 T15184	hypothetical prote
14	513	22.0	453	2 T00724	presenilin homolog
15	274	11.7	465	2 A43459	sperm membrane pro
16	119	5.1	531	2 T11596	hypothetical prote
17	110	4.7	2016	2 A38195	sodium channel pro
18	109	4.7	1840	1 CHR7M1	sodium channel pro
19	106	4.5	398	2 H75043	mg2+ transport pro
20	104	4.5	1476	1 A39901	cystic fibrosis tr
21	104	4.5	1476	1 A40303	cystic fibrosis tr
22	104	4.5	826	2 T02268	potassium transpor
23	103.5	4.4	1450	2 JC6139	cystic fibrosis tr
24	103	4.4	382	2 J47882	ubiquinol--cytochr
25	102.5	4.4	893	2 A47550	bride of sevenless
26	102	4.4	379	2 I48135	ubiquinol--cytochr
27	101.5	4.3	1681	2 A55138	sodium channel mRNA
28	100.5	4.3	1951	2 S00320	sodium channel pro
29	100.5	4.3	1983	2 A60054	isp4 protein homol
30	100.5	4.3	766	2 T01900	integrin-associate
31	100.5	4.3	324	2 S36646	chromone-c oxida
32	100	4.3	592	2 E70488	serotonin receptor
33	99.5	4.3	379	2 JC6178	sodium channel pro
34	99	4.2	2019	2 A33996	probable amino aci
35	99	4.2	521	2 T11710	

ALIGNMENTS

RESULT 1

I58098

ES-1 protein - human

C:Species: Homo sapiens (man)

C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 29-Sep-1999

C:Accession: I58098

R:Rogaev, E.I.; Sherrington, R.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Liang, Y.; C.; Cohen, D.; Lannfelt, L.; Fraser, P.E.; Rommens, J.M.; St George-Hyslop, P.H.

Nature 376, 775-778, 1995

A:Title: Familial Alzheimer's disease in kindreds with missense mutations in a gene o

A:Reference number: I58098; MUID:95379971

A:Accession: I58098

A:Status: preliminary; translated from GB/EMBL/DBDJ

A:Molecule type: mRNA

A:Residues: 1-448 <RES>

A:Cross-references: GB:L44577; NID:g950347; PIDN:AAC42012.1; PID:g950348

C:Genetics:

A:Gene: ES-1

C:Superfamily: presenilin

Query Match	100.0%;	Score	2336;	DB	2;	Length	448;		
Best Local Similarity	100.0%;	Pred. No.	8.1e-167;	Mismatches	0;	Indels	0;	Gaps	0;
1	MLTFMASDSEEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQWRSQNEEDGE	60							
Db	1	MLTFMASDSEEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQWRSQNEEDGE	60						
61	DRYVCSGVPGRPGLEELTLKYGAKHVIMLFVPTLCMIVVATIKSVRYETKNGOLI	120							
Db	61	DRYVCSGVPGRPGLEELTLKYGAKHVIMLFVPTLCMIVVATIKSVRYETKNGOLI	120						
121	YTPFTDTPSYGQRLNLSVNTLIMISIVVMTIFLVLYKYRCYKFIHGLIMSSMLL	180							
Db	121	YTPFTDTPSYGQRLNLSVNTLIMISIVVMTIFLVLYKYRCYKFIHGLIMSSMLL	180						
181	FLFTYIYLGEVLKTYNVAMDYPTLLLTWNFGAVGMCIIHWKGPLVLOQAYLIMISALMA	240							
Db	181	FLFTYIYLGEVLKTYNVAMDYPTLLLTWNFGAVGMCIIHWKGPLVLOQAYLIMISALMA	240						
241	LVFIKYLPEWSAWITLGAISYVDLVAVLCPKGPLRMLVETAQERNEPIFPALIISSAMVW	300							
Db	241	LVFIKYLPEWSAWITLGAISYVDLVAVLCPKGPLRMLVETAQERNEPIFPALIISSAMVW	300						
301	TVMKALDPPSSQALQLPYDPEMEEDSDSFGESPYEVEFPPLTGYPGELEEEERGV	360							
Db	301	TVMKALDPPSSQALQLPYDPEMEEDSDSFGESPYEVEFPPLTGYPGELEEEERGV	360						
361	KLGLGDFTFYSLVYGKAAATGSGDNTTTLACFVAILIGLCLTLLLLAVFKKALPALPISI	420							
Db	361	KLGLGDFTFYSLVYGKAAATGSGDNTTTLACFVAILIGLCLTLLLLAVFKKALPALPISI	420						
421	TFGLIFYFSTDNLVRPFMDTLASHOLYI	448							
Db	421	TFGLIFYFSTDNLVRPFMDTLASHOLYI	448						

NADH dehydrogenase
sodium channel alp
sodium channel alp
sodium channel alp
hypothetical prote
voltage-gated sodi
endothelin recepto
ubiquinol--cytochr
NADH dehydrogenase

Qy 7 SDSEEVCDERTSLMSAESPTPSRQCEGRQGPEDGENTAQRWSENEEDGEBDDPRYVCS 66
Db 5 SDSEDECNERTSLITSESPPLPSYQDGVQASEGLETSYHRERQPDSTQNNED----- 58
Qy 67 GVP-GRPGCL-----EEELTKYGAHVIMLFVPTLCMVVVATIKSVRYTT 113
Db 58 -VPNGRTSGADAYNSETTVNEEEELTKYGAHVIMLFVPTLCMVVVATIKSVRYTT 116
Qy 114 EKNGLIYTPETEDTPSYGQRLNLSVNTLMISVIVMTIFLVLYKYRCKYFHGWL 173
Db 117 EKDGOLIYTPESDTPSYGRLNLSVNTLMISVIVMTIFLVLYKYRCKYFHGWL 176
Qy 174 MSSMLLELFYIYLGELVKYNNVAMDYPTLLTWNVFAGVMGVCIHWKGPVLQOAYLI 233
Db 177 LSSMLLELFYIYLGELVKYNNVAMDYPTLLTWNVFAGVMGVCIHWKGPVLQOAYLI 236
Qy 234 MISALMALVFIKYLPEWSAWILGAISYDVLAVLCPKGPLRMLVETAQRNEPIFPALI 293
Db 237 MISALMALVFIKYLPEWSAWILGAISYDVLAVLCPKGPLRMLVETAQRNEPIFPALI 296
Qy 294 YSSAMWTVGMAKLDPPSQGAL--QLPY-----DPEMEDSDTSDFGEPSPYEPFEPPLTGY 347
Db 297 YSSAMWTVGMAKLDPPSQGAL--QLPY-----DPEMEDSDTSDFGEPSPYEPFEPPLTGY 347
Qy 348 PGELEEEERGVKLGIDGFIFYSVLVGKAAATGSGDNTTILACFVAILIGLCLTLLLA 407
Db 349 NLSEDDPEERGKVLGIDGFIFYSVLVGKAAATGSGDNTTILACFVAILIGLCLTLLLA 408
Qy 408 VFKKALPALPISITFGLIFYFSTDNLVPRFMDTLASHOLYI 448
Db 409 VFKKALPALPISITFGLIFYFSTDNLVPRFMDTLASHOLYI 449
RESULT 5
I78388
S182 protein - mouse
C:Species: Mus musculus (house mouse)
C:Date: 27-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 29-Sep-1999
C:Accession: I78388
R:Sherrington, R.; Rogava, E.I.; Liang, Y.; Rogava, E.A.; Levesque, G.; Ikeda, M.; Chiro,
E.; Pines, J.; Nee, L.; Chumakov, I.; Pollen, D.; Brookes, A.; Sanseau, P.; Pollen,
E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 375, 754-760, 1995
A:Title: Cloning of a gene bearing missense mutations in early-onset familial Alzheimer
A:Reference number: 158095; MUID:95319502
A:Accession: I78388
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-467 <RES>
A:Cross-references: GB:I42177; NID:G904129; PIDN:AAC42094.1; PID:G904130
C:Superfamily: presenilin

Query Match 62.8%; Score 1468; DB 2; Length 467;
Best Local Similarity 64.08; Pred. No. 3.9e-102;
Matches 299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;
Qy 24 ESPTPRSQCEGRQGPEDGENTAQRWSENEEDGEBDDPRYVCSGVP-----GRPPG----- 75
Db 3 EIPAPLSYFQNAQMSDSSSHSAIRSQNDSEORQOQDRQLNDPEPISNGRPSQNSRQV 62
Qy 75 -----LEEELTKYGAHVIMLFVPTLCMVVVATIKSVRYTEKNGQLIYTPFTEDT 128
Db 63 VEQDEEDELTKYGAHVIMLFVPTLCMVVVATIKSVRYTEKNGQLIYTPFTEDT 122
Qy 129 PSVGORLNSVNTLMISVIVMTIFLVLYKYRCKYFIHGWLIMSSIMLFLFYIYL 188
Db 123 ETVGORLNSVNTLMISVIVMTIFLVLYKYRCKYFIHGWLIMSSIMLFLFYIYL 182
Qy 189 GEVLTATYVNDYPTLLTWNVFAGVMGVCIHWKGPVLQOAYLIMISALMALVFIKYL 248
Db 183 GEVLTATYVNDYPTLLTWNVFAGVMGVCIHWKGPVLQOAYLIMISALMALVFIKYL 242

Qy 249 EWSAWILGAISYDVLAVLCPKGPLRMLVETAQRNEPIFPALYSSAMWTVGMAKLD 308
Db 243 EWTAWLILAVISYDVLAVLCPKGPLRMLVETAQRNEPIFPALYSSAMWTVGMAKLD 302
Qy 309 PSSQALQLPYDPEME-----EDSYDSFGEPSPYEPFEPPLTGYPG----- 350
Db 303 PEAQ--RRVPKPKYNTQRAERETQDSGSGNDGDFSEWEAQRDQSHLGRHSTPESRAA 360
Qy 350 -EEL-----EEERGVKLGIDGFIFYSVLVGKAAATGSGDNTTILACFVAILIGLCL 401
Db 361 VOELSGSILTSDDPEERGKVLGIDGFIFYSVLVGKAAATGSGDNTTILACFVAILIGLCL 420
Qy 402 TLLLAVERKALPALPISITFGLIFYFSTDNLVPRFMDTLASHOLYI 448
Db 421 TLLLAVERKALPALPISITFGLIFYFSTDNLVPRFMDTLASHOLYI 467
RESULT 6
S58396
Presenilin 1, splice form 467 - human
N:Alternate names: Alzheimer's disease protein 3; protein S182
C:Species: Homo sapiens (man)
C:Date: 29-Jan-1998 #sequence_revision 13-Feb-1998 #text_change 29-Sep-1999
C:Accession: S58396; S71401; S71402
R:Sherrington, R.; Rogava, E.I.; Liang, Y.; Rogava, E.A.; Levesque, G.; Ikeda, M.; Chiro,
E.; Pines, J.; Nee, L.; Chumakov, I.; Pollen, D.; Brookes, A.; Sanseau, P.; Pollen,
E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 375, 754-760, 1995
A:Title: Cloning of a gene bearing missense mutations in early-onset familial Alzheimer
A:Reference number: 158095; MUID:95319502
A:Accession: S58396
A:Molecule type: mRNA
A:Residues: 1-467 <SHE>
A:Cross-references: EMBL:I42110; NID:G904118; PIDN:AAB46416.1; PID:G904119
R:Vidal, R.; Ghiso, J.; Wisniewski, T.; Frangione, B.
FEBS Lett. 393, 19-23, 1996
A:Title: Alzheimer's presenilin 1 gene expression in platelets and megakaryocytes. Id
A:Reference number: S71401; MUID:96397521
A:Accession: S71401
A:Status: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 24-32;254-256,290-292;316-317,376-379 <VIW>
A:Experimental source: Dani megakaryotic cell line (ATCC CRL-9792) and platelets
C:Genetics:
A:Gene: GDB:PSEN1; AD3; PAD; S182; PS1
A:Cross-references: GDB:I35682; OMIM:104311
A:Map position: 14q24.3-14q24.3
C:Superfamily: presenilin
C:Keywords: alternative splicing; Alzheimer's disease; glycoprotein; transmembrane pr
F:82-100/Domain: transmembrane #status predicted <TM1>
F:133-154/Domain: transmembrane #status predicted <TM2>
F:164-185/Domain: transmembrane #status predicted <TM3>
F:195-213/Domain: transmembrane #status predicted <TM4>
F:221-238/Domain: transmembrane #status predicted <TM5>
F:244-264/Domain: transmembrane #status predicted <TM6>
F:281-301/Domain: transmembrane #status predicted <TM7>
F:408-428/Domain: transmembrane #status predicted <TM8>
F:433-453/Domain: transmembrane #status predicted <TM9>
F:279,405/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 62.8%; Score 1467; DB 2; Length 467;
Best Local Similarity 65.7%; Pred. No. 4.6e-102;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;
Qy 24 ESPTPRSQCEGRQGPEDGENTAQRWSENEEDGEBDDPRYVCSGVP-----GRPPG----- 75
Db 3 EIPAPLSYFQNAQMSDSSSHSAIRSQNDSEORQOQDRQLNDPEPISNGRPSQNSRQV 61
Qy 75 -----LEEELTKYGAHVIMLFVPTLCMVVVATIKSVRYTEKNGQLIYTPFTED 127
Db 62 VVEQDEEDELTKYGAHVIMLFVPTLCMVVVATIKSVRYTEKNGQLIYTPFTED 121

presenilin 1 protein isoform 467 - lesser mouse lemur
C:Species: Microcebus murinus (lesser mouse lemur)
C:Date: 31-Jan-1997 #sequence_revision 31-Jan-1997 #text_change 29-Sep-1999
C:Accession: JC5080
R:Calenda, A.; Mestre-Frances, N.; Czech, C.; Pradier, L.; Petter, A.; Bons, N.; Bellis,
Biochem. Biophys. Res. Commun. 228, 430-439, 1996
A:Title: Molecular cloning, sequencing, and brain expression of the presenilin 1 gene in
A:Reference number: JC5080; MUID:97079199
A:Accession: JC5080
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-467 <CAL>
A:Cross-references: EMBL:Z71333
C:Comment: This protein is an integral membrane protein with seven transmembrane domain
C:Genetics:
A:Gene: psi
A:Map position: 14
C:Superfamily: presenilin
C:Keywords: transmembrane protein
F:82-100/Domain: transmembrane #status predicted <TM1>
F:133-154/Domain: transmembrane #status predicted <TM2>
F:164-185/Domain: transmembrane #status predicted <TM3>
F:195-213/Domain: transmembrane #status predicted <TM4>
F:221-238/Domain: transmembrane #status predicted <TM5>
F:244-261/Domain: transmembrane #status predicted <TM6>
F:408-428/Domain: transmembrane #status predicted <TM7>

Query Match 61.9%; Score 1447; DB 2; Length 467;
Best Local Similarity 64.4%; Pred. No. 1.4e-100;
Matches 300; Conservative 41; Mismatches 83; Indels 42; Gaps 7;
Qy 24 ESPTPRSCQEGRGQPEDGENTAQWRSQNEEDGDDPRYVCSGVVP-----GRPPG----- 75
Db 3 ELPAPLSYFQNAQMSDNLNSTRQNDREQDGHDRRL-GNPELSNGRPOGNSGP 61

Qy 75 -----LEEELTKYGAHVIMLFVPTLCMVVVVATIKSVFRFTKNGLIYTPFTED 127
Db 62 VVERDEEDELTKYGAHVIMLFVPTLCMVVVVATIKSVFRFTKNGLIYTPFTED 121

Qy 128 TPSVGORLLNSVNTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSMLLFLFTYI 187
Db 122 TETVQGRALSHVLAAMISVIVMTIFLVLYKYRCYKFIHAWLIISLLLLFFFSFI 181

Qy 188 LGEVLKTYNVAMDPTLLTWNFGAVGVCIHMKGPLVLAQAYLIMISALMALVFIKYL 247
Db 182 LGEVFTYNVAVDYITVALLIWNFGVGMISIHMKGPLRLQAYLIMISALMALVFIKYL 241

Qy 248 PEWSAWVLGAISYVDLVAVLCPLKPLRMVETAGERNPIFPALIIYSAMVWTVGMAKL 307
Db 242 PEWTAWLLAVISYVDLVAVLCPLKPLRMVETAGERNETLFPALIIYSSTWVLYNMAEG 301

Qy 308 DPSSQGLAL--QLPYD-----PEMEEDSYDSFGSPSYPEVFP-----PLTGYPG 349
Db 302 DPEAQRVSKNTKYNAQGTREAAQSVPEDDGGFSEWEAQRDSQLGPHRSTSVSRAAV 361

Qy 350 EEL-----EEEEERGKVLGIDGFIFYSVLVGKAAATGSGDNTTTLACFVAILGLCLT 402
Db 362 QEISSIPASEDPEERGKVLGIDGFIFYSVLVGKASATASGDWNTTIACFVAILGLCLT 421

Qy 403 LLLAVFKKALPALPISITFGLIFFYFSDNLVVRPMDTLASHQLYI 448
Db 422 LLLAIFKKALPALPISITFGLVFFATDYLVPMDQLAFHOFYI 467

RESULT 10
JC5081
presenilin 1 protein isoform 463 - lesser mouse lemur
C:Species: Microcebus murinus (lesser mouse lemur)
C:Date: 31-Jan-1997 #sequence_revision 31-Jan-1997 #text_change 13-Sep-1998
C:Accession: JC5081
R:Calenda, A.; Mestre-Frances, N.; Czech, C.; Pradier, L.; Petter, A.; Bons, N.; Bellis,

Biochem. Biophys. Res. Commun. 228, 430-439, 1996
A:Title: Molecular cloning, sequencing, and brain expression of the presenilin 1 gene
A:Reference number: JC5080; MUID:97079199
A:Accession: JC5081
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-463 <CAL>
A:Cross-references: EMBL:Z71333
C:Comment: This protein is an intermembrane protein with seven transmembrane domains.
C:Genetics:
A:Gene: psi
A:Map position: 14
C:Superfamily: presenilin
C:Keywords: transmembrane protein
F:78-96/Domain: transmembrane #status predicted <TM1>
F:129-150/Domain: transmembrane #status predicted <TM2>
F:160-181/Domain: transmembrane #status predicted <TM3>
F:191-209/Domain: transmembrane #status predicted <TM4>
F:217-234/Domain: transmembrane #status predicted <TM5>
F:240-257/Domain: transmembrane #status predicted <TM6>
F:404-424/Domain: transmembrane #status predicted <TM7>

Query Match 61.6%; Score 1440; DB 2; Length 463;
Best Local Similarity 64.2%; Pred. No. 4.7e-100;
Matches 300; Conservative 39; Mismatches 80; Indels 48; Gaps 8;
Qy 24 ESPTPRSCQEGRGQPEDG--ENTAQWRSQNEED-----GEEDPRYVCSGVPPRG-- 75
Db 3 ELPAPLSYFQNAQMSDNLNSTRQNDREQDGHDRRLGNPELS-----NGRPOGNSG 56

Qy 75 -----LEEELTKYGAHVIMLFVPTLCMVVVVATIKSVFRFTKNGLIYTPFTE 126
Db 57 PVVERDEEDELTKYGAHVIMLFVPTLCMVVVVATIKSVFRTRKDGQLIYTPFTE 116

Qy 127 DPSPVGORLLNSVNTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSMLLFLFTYI 186
Db 117 DPTVQGRALSHVLAAMISVIVMTIFLVLYKYRCYKFIHAWLIISLLLLFFFSFI 176

Qy 187 YLGEVLTNYNVAMDPTLLTWNFGAVGVCIHMKGPLVLAQAYLIMISALMALVFIKY 246
Db 177 YLGEVFTYNVAVDYITVALLIWNFGVGMISIHMKGPLRLQAYLIMISALMALVFIKY 236

Qy 247 PEWSAWVLGAISYVDLVAVLCPLKPLRMVETAGERNPIFPALIIYSAMVWTVGMAK 306
Db 237 LPEWTAWLLAVISYVDLVAVLCPLKPLRMVETAGERNETLFPALIIYSSTWVLYNMAE 296

Qy 307 LDPSSQGLAL--QLPYD-----PEMEEDSYDSFGSPSYPEVFP-----PLTGYP 348
Db 297 GDPEAQRVSKNTKYNAQGTREAAQSVPEDDGGFSEWEAQRDSQLGPHRSTSVSRAA 356

Qy 349 GEEL-----EEEEERGKVLGIDGFIFYSVLVGKAAATGSGDNTTTLACFVAILGLCL 401
Db 357 VQEISSIPASEDPEERGKVLGIDGFIFYSVLVGKASATASGDWNTTIACFVAILGLCL 416

Qy 402 TLLLAIFKKALPALPISITFGLIFFYFSDNLVVRPMDTLASHQLYI 448
Db 417 TLLLAIFKKALPALPISITFGLVFFATDYLVPMDQLAFHOFYI 463

RESULT 11
S63684
presenilin 1, splice form 374 - human
N:Alternate names: Alzheimer's disease protein 3
C:Species: Homo sapiens (man)
C:Date: 20-Jul-1996 #sequence_revision 13-Mar-1997 #text_change 29-Sep-1999
C:Accession: S63684
R:Sahara, N.; Yahagi, Y.; Takagi, H.; Kondo, T.; Okochi, M.; Usami, M.; Shirasawa, T.
FEBS Lett. 381, 7-11, 1996
A:Title: Identification and characterization of presenilin I-467, I-463 and I-374.
A:Reference number: S63683; MUID:96193901
A:Accession: S63684

||||| : || | | : |
Db 336 YFSSHALPFTDLCTSQLILI 358

RESULT 14

T00724

presenilin homolog F22013.19 - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 12-Feb-1999 #sequence_revision 12-Feb-1999 #text_change 20-Sep-1999
C:Accession: T00724
R:Shinn, P.; Buehler, E.; Dewar, K.; Peng, J.; Kim, C.; Li, Y.; Sun, H.; Conway, A.; Con-
eologis, A.; Ecker, J.R.
submitted to the EMBL Data Library, April 1998
A:Description: Genomic sequence for Arabidopsis thaliana BAC F22013.
A:Reference number: Z14200
A:Accession: T00724
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-453 <SHI>
A:Cross-references: EMBL:AC003981; NID:g3063438; PID:g3063457; GSPDB:GNO059; ATSP:F22013
C:Genetics:
A:Gene: ATSP:F22013.19
A:Map position: 1
A:Introns: 108/1
C:Superfamily: presenilin

Query Match 22.08; Score 513; DB 2; Length 453;

Best Local Similarity 31.18; Pred. No. 5.2e-31;

Matches 136; Conservative 78; Mismatches 130; Indels 94; Gaps 12;

QY 88 VIMLFVPTLCMIVVATIKSVREYTEKN----GQLIYPTFTEDTPSVGQRLLSVNLTL 143
Db 13 IIGVMPVSCMFLVLLTSLVTSQPKIRSAANLIYIENPDSSTTV--KLEGLANAI 70
QY 144 IMISVIVMTIFVLVKYRCYKFIHGLWIMSLMLLFLEFYIYVLGEVLKTYNVAMDYPT 203
Db 71 VFVVLIAAVTIFVLVLYYNTFNELKHYMRSAFFVLGTMGGAIFLSIIQHFSIPVDSIT 130
QY 204 LLTVNFWGAVGVCIIHWK--PLVLOQAYLIMISALMALVFIKYLPEWSAWILGAISVY 262
Db 131 CFILLNFITLIGLSVFAGGPIVLRQCYMVGMIVVAAWFK-LPEWTTWFIIVALALY 189
QY 263 DLVAVLCPKPLMLVETAQERNEPIPALIY-----SSAMVTVGMAKL 307
Db 190 DLVAVLAPGGLKLLVYELASRDEEL-PAMYEARPTVSSGNQRNRGSSRLALVGGGV 248
QY 308 DPSOGALQLPYDEM----EEDSY-----DSFGPSYPPEVPEPLTGY 347
Db 249 SDGSEVLOAVRHDVNLGRENHNDYNAIVRDIDNVDDGIGNGSRGGLERSPLVGS 308
QY 348 PG-----LEEEE-----EE-----ERGV 360
Db 309 PSASEHSTVGTGRNMDRESVDEMSPLVELMGWGNREARGLESNDNVDSINRGI 368
QY 361 KLGLGDFIFYSVLGKAAATGSGDWTTLACFVAILIGLCTLLALLAVFKKALPALPISI 420
Db 369 KLGLGDFIFYSVLGKAAATGSGDWTTLACFVAILIGLCTLLALLAVFKKALPALPISI 420
QY 421 TFLGLIFYFTDNLVRPFM 438
Db 426 MGVVFFLRLMEPFV 443

RESULT 15

A43459

sperm membrane protein spe-4 - Caenorhabditis elegans
N:Alternate names: probable integral membrane protein
C:Species: Caenorhabditis elegans
C:Date: 10-Jun-1993 #sequence_revision 18-Nov-1994 #text_change 09-Sep-1997
C:Accession: A43459; S24632; S24633
R:L'Hernault, S.W.; Arduengo, P.M.
J. Cell Biol. 119, 55-68, 1992

A:Title: Mutation of a putative sperm membrane protein in *Caenorhabditis elegans* prev
A:Reference number: A43459; MUID:92407040
A:Accession: A43459
A:Status: preliminary; not compared with conceptual translation
A:Molecule type: DNA; mRNA
A:Residues: 1-465 <LHE>
A:Cross-references: EMBL:Z14067; NID:g6868; PID:g6869; EMBL:Z14066; NID:g6870; PID:g6
A:Experimental source: strain Bristol N2
A:Note: the nucleotide sequence was submitted to the EMBL Data Library, July 1992
C:Genetics:
A:Introns: 69/3; 154/3; 200/1; 224/3; 300/1; 386/1; 435/1

Query Match 11.78; Score 274; DB 2; Length 465;

Best Local Similarity 21.58; Pred. No. 3.4e-13;

Matches 93; Conservative 80; Mismatches 127; Indels 132; Gaps 14;

QY 114 EKNGLIYPTFTEDT--PSVGORLLNSVLN----LFTYIYVLGEVLKTYNVAMDYPTLL-----TVNFGA 213
Db 42 EVNSLSKTYFLDPSFEQTGNLLLDGFIINGVGTILVIGCVSFIMLAF--VLEDFR--RI 97
QY 168 IHGWLIMSLMLLF-----LFTYIYVLGEVLKTYNVAMDYPTLL-----TVNFGA 213
Db 98 VKAWLTLSCLLILFGVSAQTLHDMFSQVFDQDDNQY-----YMTIVLIWVTVVYVGF- 152
QY 214 VGVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWILGAISVYDLVAVLCPKGP 273
Db 152 -GIYAFFSNSLLIHLQIFVVTNCSLISVFLRVFPFSKTTFWLWILVFLWDLFAVLAPMG 210
QY 274 LRMLVETAQERNEPIPALIYSSAMVTVGMAKLDPSSOG-----ALQLPYDPE 322
Db 211 LKKVQERASDYKCVLNLMFSANEKRLTAGSNQETNEGESTIRRTVKQIIEYTKRE 270
QY 323 MEEDSY-----DSP-----GEPSPYEPPEPLTGYGPEEEEEEE--- 358
Db 271 AQDDEFYQKIRQRAAINPDSVPTESPLVEAEPSPIELKEKNST----EELSDDSDTS 326
QY 358 -----R 358
Db 327 ETSGSSSLSSSDSTTVSTSDISTAECDQKEDDLVSNLSPNNDKRPATAADALNDGE 386
QY 359 GVKLGLGDFIFYSVLGKAAATGSGDWTTLACFVAILIGLCTLLALLAVFKKALPALPI 418
Db 387 VRLUGFGDFVYFSLIGQAASGCP--FAVISAALGILFGLVVLTVFVSTEESTTPALPL 444
QY 419 SITFGLIIFYFT 430
Db 445 PVICGTCFCYFS 456

Search completed: March 18, 2000, 14:11:52

Job time: 489 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 16:08:11 ; Search time 32.57 Seconds
(without alignments)
410.791 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFMADSDSEEVCDERTSL.....STDNLVRFPMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 82229 seqs, 29864866 residues

Database : SwissProt_38:*

Word size : 0

Number of hits that pass the threshold : 82229

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	2336	100.0	448	1	PSN2_HUMAN
2	2232	95.5	448	1	PSN2_MOUSE
3	2227	95.3	448	1	PSN2_RAT
4	2218	94.9	445	1	PSN2_MICMU
5	1624.5	69.5	449	1	PSN2_XENLA
6	1468	62.8	467	1	PSN1_MOUSE
7	1467	62.8	467	1	PSN1_HUMAN
8	1459.5	62.5	468	1	PSN1_RAT
9	1449.5	62.1	433	1	PSN1_XENLA
10	1447	61.9	467	1	PSN1_MICMU
11	1150	49.2	541	1	PSN1_DROME
12	1043	44.6	836	1	YLAK_CAEEL
13	983	42.1	461	1	SE12_CAEEL
14	524.5	22.5	358	1	HOP1_CAEEL
15	513	22.0	453	1	PSNH_ARATH
16	274	11.7	465	1	SPEX_CAEEL
17	119	5.1	531	1	YDFG_SCHPO
18	110	4.7	2016	1	CIN5_HUMAN
19	109	4.7	1840	1	CIN4_RAT
20	108	4.6	381	1	CYB_DASCR
21	104	4.5	1476	1	CFTF_MOUSE
22	103.5	4.4	1450	1	CFTF_RABIT
23	103	4.4	382	1	CYB_DIDMA
24	103	4.4	380	1	CYB_MICLO
25	102.5	4.4	893	1	BOSS_DROVI
26	102.5	4.4	381	1	CYB_ANTIPL
27	102	4.4	381	1	CYB_DASMA
28	102	4.4	381	1	CYB_PSENI
29	101.5	4.3	381	1	CYB_NINIV
30	101.5	4.3	381	1	CYB_PANAP
31	101	4.3	381	1	CYB_PLAMS
32	100.5	4.3	1951	1	CIN3_RAT
33	100.5	4.3	381	1	CYB_PSEMD
34	100	4.3	381	1	CYB_DASGE
35	99	4.2	2019	1	CIN5_RAT
36	99	4.2	381	1	CYB_ANTI
37	99	4.2	381	1	CYB_PHATA
38	98.5	4.2	872	1	CIN3_HUMAN
39	98	4.2	1836	1	CIN4_HUMAN

40	98	4.2	381	1	CYB_ANTI	Q33782 antechinus
41	98	4.2	381	1	CYB_DASAL	Q34289 dasyurus al
42	98	4.2	381	1	CYB_SMICR	Q35810 smnthopsis
43	98	4.2	460	1	NU4M_GADMO	P55781 gadus morhu
44	98	4.2	217	1	YPRA_ECOLI	P13974 escherichia
45	97.5	4.2	381	1	CYB_DASHA	Q34321 dasyurus ha

ALIGNMENTS

RESULT 1						
PSN2_HUMAN						
ID PSN2_HUMAN	STANDARD;	PRT;	448 AA.			
AC P49810;						
DT 01-OCT-1996	(Rel. 34, Created)					
DT 01-OCT-1996	(Rel. 34, Last sequence update)					
DT 15-JUL-1999	(Rel. 38, Last annotation update)					
DE PRESENTLIN 2 (PS-2) (STM-2) (E5-1) (AD3LP) (AD5).						
GN PSN2 OR PSN12 OR AD4 OR PS2 OR STM2.						
OS Homo sapiens (Human).						
OC Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;						
OC Eutheria; Primates; Catarrhini; Homiidae; Homo.						
RN [1]						
RP SEQUENCE FROM N.A., AND VARIANT FAD ILE-141.						
RX MEDLINE; 95365816.						
RA LEVY-LAHAD E., WASCO W., POORRAJ P., ROMANO D.M., OSHIMA J.,						
RA PETTINGELL W.H., YU C.-E., JONDRO P.D., SCHMIDT S.D., WANG K.,						
RA CROWLEY A.C., FU Y.-H., GUENETTE S.Y., GALAS D., NEMENS E.,						
RA WIJSMAN E.M., BIRD T.D., SCHELLENBERG G.D., TANZI R.E.;						
RT "Candidate gene for the chromosome 1 familial Alzheimer's disease						
RT locus.";						
RL Science 269:973-977(1995).						
RN [2]						
RP SEQUENCE FROM N.A., AND VARIANTS FAD ILE-141 AND VAL-239.						
RC TISSUE-BRAIN, AND COLON;						
RX MEDLINE; 95379971.						
RA ROGAEV E.I., SHERRINGTON R., ROGAEVA E.A., LEVESQUE G., IKEDA M.,						
RA LIANG Y., CHI H., LIN C., HOLMAN K., TSUDA T., MAR L., SORBI S.,						
RA NACMIAS B., PIACENTINI S., AMADUCCI L., CHUMAKOV I., COHEN D.,						
RA LANFELT L., FRASER P.E., ROMMENS J.M., ST GEORGE-HYSLOP P.H.;						
RT "Familial Alzheimer's disease in kindreds with missense mutations in						
RT a gene on chromosome 1 related to the Alzheimer's disease type 3						
RT gene.";						
RL Nature 376:775-778(1995).						
RN [3]						
RP SEQUENCE FROM N.A.						
RX MEDLINE; 96109229.						
RA LI J., MA J., POTTER H.;						
RT "Identification and expression analysis of a potential familial						
RT Alzheimer disease gene on chromosome 1 related to AD3.";						
RL Proc. Natl. Acad. Sci. U.S.A. 92:12180-12184(1995).						
RN [4]						
RP SEQUENCE FROM N.A.						
RA LEVY-LAHAD E., POORRAJ P., WANG K., FU Y.H., OSHIMA J.,						
RA MULLIGAN J., SCHELLENBERG G.D.;						
RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.						
RN [5]						
RP REVIEW ON VARIANTS.						
RX MEDLINE; 98180715.						
RA CRUTS M., VAN BROECKHOVEN C.;						
RT "Presenilin mutations in Alzheimer's disease.";						
RL Hum. Mutat. 11:183-190(1998).						
RN [6]						
RP VARIANT AD HIS-62.						
RX MEDLINE; 98046005.						
RA CRUTS M., VAN DUJN C.M., BACKHOVENS H., VAN DEN BROECK M.,						
RA WEHNET A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J.,						
RA ST GEORGE-HYSLOP P.H., HOFMAN A., VAN BROECKHOVEN C.;						
RT "Estimation of the genetic contribution of presenilin-1 and -2						
RT mutations in a population-based study of presenile Alzheimer						
RT disease.";						
RL Hum. Mol. Genet. 7:43-51(1998).						

DR EMBL; U57325; AAC53311.1; -;
DR EMBL; AF038935; AAB92660.1; -;
DR EMBL; U49111; AAC52935.1; -;
DR MGD; MGI:109284; PSEN2.
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Alternative initiation.
FT CHAIN 1 448 PRESENILIN 2.
FT CHAIN 298 448 PRESENILIN 2-SHORT.
FT TRANSMEM 88 106 POTENTIAL.
FT TRANSMEM 142 160 POTENTIAL.
FT TRANSMEM 167 188 POTENTIAL.
FT TRANSMEM 203 219 POTENTIAL.
FT TRANSMEM 230 246 POTENTIAL.
FT TRANSMEM 253 269 POTENTIAL.
FT TRANSMEM 288 305 POTENTIAL.
FT TRANSMEM 387 406 POTENTIAL.
FT TRANSMEM 413 429 POTENTIAL.
FT CONFLICT 87 87 R -> H (IN REF. 2).
FT CONFLICT 226 226 A -> V (IN REF. 2).
FT CONFLICT 324 324 MISSING (IN REF. 2).
SQ SEQUENCE 448 AA; 49955 MW; 680ACF19 CRC32;

Query Match 95.5%; Score 2232; DB 1; Length 448;
Best Local Similarity 95.5%; Pred. No. 7.3e-147;
Matches 428; Conservative 6; Mismatches 14; Indels 0; Gaps 0;

QY 1 MLTFMSDSEEEVCDERTSLMSAESPTRSCQGRGQPEDGENTAQRSENEDEEDP 60
Db 1 MLAFMSDSEEEVCDERTSLMSAESPTRSCQGRGQPEDGENTAQRSENEDEEDP 60
QY 61 DRYVCSGVPGRPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVFYTEKNGQLI 120
Db 61 DRYACSGAPGRPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVFYTEKNGQLI 120
QY 121 YTPFTEDTPSGORLLNSVNTLIMISIVVMTFLVLYKYRCYKFIHGLWLSMLL 180
Db 121 YTPFTEDTPSGORLLNSVNTLIMISIVVMTFLVLYKYRCYKFIHGLWLSMLL 180
QY 181 FLFTYIYLGEVLKTYNNVAMDYPTLLLVNMFAGVMYCIHWKGPLVLOQAYLIVISALMA 240
Db 181 FLFTYIYLGEVLKTYNNVAMDYPTLLLVNMFAGVMYCIHWKGPLVLOQAYLIVISALMA 240
QY 241 LVFIKILPEWSANVILGAISYDVLVAVLCRGPPLMLVETQAQERNEIFPALIYSSAMVW 300
Db 241 LVFIKILPEWSANVILGAISYDVLVAVLCRGPPLMLVETQAQERNEIFPALIYSSAMVW 300
QY 301 TVGMAKLDPSQCALQLPYPMEEDSYDSFGSPSYPEVEPPLTGYPGRELEEEERGV 360
Db 301 TVGMAKLDPSQCALQLPYPMEEDSYDSFGSPSYPEVEPPLTGYPGRELEEEERGV 360
QY 361 KLGLGDFIFYSVLVGAAATGSDGWNNTLACFAVAILGLCITLLLLAVFKKALPALPISI 420
Db 361 KLGLGDFIFYSVLVGAAATGSDGWNNTLACFAVAILGLCITLLLLAVFKKALPALPISI 420
QY 421 TFGLIIFYSTDLNLRPMDTFLASHQLYI 448
Db 421 TFGLIIFYSTDLNLRPMDTFLASHQLYI 448

RESULT 3
PSN2_RAT
ID PSN2_RAT STANDARD; PRT; 448 AA.
AC O88777; O35546; O08947;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN 2 (PS-2).
GN PSN2 OR PSN2 OR PS2.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]

RP SEQUENCE FROM N.A.
RC STRAIN-WISTAR; TISSUE-BRAIN;
RA FRENZEL S., ABDEL A.S., LUEBBERT H.;
RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-WISTAR; TISSUE-BRAIN;
RX MEDLINE; 97473536
RA TAKAHASHI H., MERCKEN M., NAKAZATO Y., NOGUCHI K., MURAYAMA M.,
RA IMAHORI K., TAKASHIMA A.;
RT "Cloning of CDNA and expression of the gene encoding rat
presenilin-2";
RL Gene 197:383-387(1997).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN-WISTAR; TISSUE-BRAIN;
RX MEDLINE; 98207716
RA TANAHASHI H., TABIRA T.;
RT "Cloning of the CDNA encoding rat presenilin-2";
RL Biochim. Biophys. Acta 1396:259-262(1998).
CC -!- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE
CC EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE. MAY
CC FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS (BY
CC SIMILARITY).
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC
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DR EMBL; X99267; CAA67663.1; -;
DR EMBL; D83700; BAA22832.1; -;
DR EMBL; AB004454; BAA20406.1; -;
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane.
FT TRANSMEM 88 106 POTENTIAL.
FT TRANSMEM 142 160 POTENTIAL.
FT TRANSMEM 167 188 POTENTIAL.
FT TRANSMEM 203 219 POTENTIAL.
FT TRANSMEM 230 246 POTENTIAL.
FT TRANSMEM 253 269 POTENTIAL.
FT TRANSMEM 288 305 POTENTIAL.
FT TRANSMEM 387 406 POTENTIAL.
FT TRANSMEM 413 429 POTENTIAL.
FT CONFLICT 7 7 S -> T (IN REF. 1).
FT CONFLICT 86 87 KH -> ND (IN REF. 3).
SQ SEQUENCE 448 AA; 50051 MW; E80DF681 CRC32;

Query Match 95.3%; Score 2227; DB 1; Length 448;
Best Local Similarity 95.3%; Pred. No. 1.6e-146;
Matches 427; Conservative 6; Mismatches 15; Indels 0; Gaps 0;

QY 1 MLTFMSDSEEEVCDERTSLMSAESPTRSCQGRGQPEDGENTAQRSENEDEEDP 60
Db 1 MLTFMSDSEEEVCDERTSLMSAESPTRSCQGRGQPEDGENTAQRSENEDEEDP 60
QY 61 DRYVCSGVPGRPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVFYTEKNGQLI 120
Db 61 DRYACSGVPGRPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVFYTEKNGQLI 120
QY 121 YTPFTEDTPSGORLLNSVNTLIMISIVVMTFLVLYKYRCYKFIHGLWLSMLL 180
Db 121 YTPFTEDTPSGORLLNSVNTLIMISIVVMTFLVLYKYRCYKFIHGLWLSMLL 180
QY 181 FLFTYIYLGEVLKTYNNVAMDYPTLLLVNMFAGVMYCIHWKGPLVLOQAYLIVISALMA 240
Db 181 FLFTYIYLGEVLKTYNNVAMDYPTLLLVNMFAGVMYCIHWKGPLVLOQAYLIVISALMA 240

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FT TRANSMEM 415 435 POTENTIAL.
FT CARBOHYD 387 387 POTENTIAL.
SQ SEQUENCE 449 AA; 50274 MW; DF6EAF4E CRC32;

Query Match 69.5%; Score 1624.5; DB 1; Length 449;
Best Local Similarity 72.0%; Pred. No. 4.9e-105;
Matches 332; Conservative 34; Mismatches 60; Indels 35; Gaps 7;

Qy 7 SDSEEVCDERTSLMSASPTPRSCQGRQGPEDGENTAQRWSQNEDEGDDPDRYVCS 66
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Dy 5 SDSEEDENERTSLTSESPLPSYQDQVQASEGLETSHRERQDPSQNNED----- 58
Qy 67 GVP--GRPPGL-----EEELTKYGAKHVIMLFVPTLGMIVVWATIKSVREVT 113
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 58 -VPGRTSGADAYNSETTVNEEEELTKYGARHVIMLFVPTLGMIVVWATIKSVREVT 116
Qy 114 EKNGLIYPTFEDTPSYGQRLNSVLTLMISIVVMTIFLVVLYKYRCYKFIHGWL 173
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 117 EKDGQIYTPSEDTTSVGERLLNSVLTLMISIVVMTIFLVVLYKYRCYKFIHGWL 176
Qy 174 MSSLMFLFYIYLGELVKIYNVAMDPTLLLTWNFGAVGMCVCIHWKGPLLQQAAYLI 233
Dy :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 177 LSLEMLFEMFTYIYLVSEFKYNTIAMDPTLLFMVWTFNFGAVGMCVCIHWKGPLLQQAAYLI 236
Qy 234 MISALMALVETIKYLPWESAIVILGAISYDVLVAVLCPKGPLRMLVETAQERNEPIFPALI 293
Dy :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 237 MISALMALVETIKYLPWESAIVILGAISYDVLVAVLCPKGPLRMLVETAQERNEPIFPALI 296
Qy 294 YSSAMVTVTGMKLDPPSSOGAL--QLPY-----DPMEEDSDYDSFGESPYVEFPEPLTGY 347
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 297 YSSAMVTVTGMKLDPPSSOGAL--QLPY-----DPMEEDSDYDSFGESPYVEFPEPLTGY 347
Qy 348 PGEELBEERGVKLGIDGDFIFYSVLVGAATAATGSDWNTTILACFVAILGLTLLLA 407
Dy :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 349 NLSSDDPDEERGVLGLGDFIFYSVLVGAATAATGSDWNTTILACFVAILGLTLLLA 408
Qy 408 VFKALPALPISITFGLIFYSTDNLVRPFMDTLASHOLYI 448
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 409 VFKALPALPISITFGLIFYSTDNLVRPFMDTLASHOLYI 449

RESULT 6
PSN1_MOUSE STANDARD: PRT; 467 AA.
AC P49769;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE PRESENILIN 1 (PS-1) (S182 PROTEIN).
GN PSN1 OR PSN1 OR AD3H.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-BRAIN;
RX MEDLINE; 95319502.
RA SHERINGTON R., ROGAEV E.I., LIANG Y., ROGAEVA E.A., LEVESQUE G.,
RA IKEDA M., CHI H., LIN C., LI G., HOLMAN K., TSUDA T., MAR L.,
RA FONCIN J.-F., BRUNI A.C., MONTESI M.P., SORBI S., RAINERO I.,
RA PINESSI L., NEE L., CHUMAKOV I., POLLEN D., BROOKES A.,
RA SANSEAU P., POLINSKY R.J., WASCO W., DA SILVA H.A.R., HAINES J.L.,
RA PERICAK-VANCE M.A., TANZI R.E., ROSES A.D., FRASER P.E.,
RA ROMMENS J.M., ST GEORGE-HYSLOP P.H.;
RT "Cloning of a gene bearing missense mutations in early-onset familial
RT Alzheimer's disease."
RL Nature 375:754-760(1995).
[2]
RN SEQUENCE FROM N.A.
RP STRAIN=129/SVJ;
RX MEDLINE; 97442406.
RA MITSUDA N., ROSES A.D., VITEK M.P.;

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RT "Transcriptional regulation of the mouse presenilin-1 gene.";
RL J. Biol. Chem. 272:23489-23497(1997).
CC -!- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE
CC EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE.
CC -!- MAY FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS.
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
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CC -----
DR EMBL; L42177; AAC42094.1; -.
DR EMBL; AF007560; AAB72049.1; -.
DR MGD; MGI:1202717; PSEN1.
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Glycoprotein.
FT TRANSMEM 83 103 POTENTIAL.
FT TRANSMEM 133 153 POTENTIAL.
FT TRANSMEM 161 181 POTENTIAL.
FT TRANSMEM 195 215 POTENTIAL.
FT TRANSMEM 221 241 POTENTIAL.
FT TRANSMEM 241 264 POTENTIAL.
FT TRANSMEM 281 301 POTENTIAL.
FT TRANSMEM 408 428 POTENTIAL.
FT TRANSMEM 433 453 POTENTIAL.
FT CARBOHYD 30 30 POTENTIAL.
FT CARBOHYD 279 279 POTENTIAL.
FT CARBOHYD 405 405 POTENTIAL.
SQ SEQUENCE 467 AA; 52639 MW; 3AE1350D CRC32;

Query Match 62.8%; Score 1468; DB 1; Length 467;
Best Local Similarity 64.0%; Pred. No. 3e-94;
Matches 299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;

Qy 24 ESPTPRSCQGRQGPEDGENTAQRWSQNEDEGDDPDRYVCSGVP-----GRPPG----- 75
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 3 EIPAPLSYFQNAQMSDESHSSAISRQNSQDSQROQHQRDLNPEPISNGRPSQNSQV 62
Qy 75 -----LEEELTKYGAKHVIMLFVPTLGMIVVWATIKSVRFYTEKNGQLIYTPFTE 128
Dy :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 63 VEQDEEDELTKYGAKHVIMLFVPTLGMIVVWATIKSVFYTRKQGLIYTPFTE 122
Qy 129 PSVGORLLNSVLTLMISIVVMTIFLVVLYKYRCYKFIHGWLIMSSLMFLFYIYL 188
Dy :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 123 ETVGORALHSILNAAIMISVIVMTILLVLYKYRCYKVIHAWLISSLLFFSFYIL 182
Qy 189 GEVLKTYNVAMDYPTLLLTWNFGAVGMCVCIHWKGPLVLOQAALIMISALMALVETIKYLP 248
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 183 GEVFTYNAVVDYTVALLIWNFGVGMIAHWKGPLRLQQAALIMISALMALVETIKYLP 242
Qy 249 EWSAWVILGAISYDVLVAVLCPKGPLRMLVETAQERNEPIFPALIYSSAMVTVTGMKLD 308
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 243 EMTAWLILAVISYDVLVAVLCPKGPLRMLVETAQERNEPIFPALIYSSAMVTVTGMKLD 302
Qy 309 PSSQALQLPYDPEME-----EDSDYDSFGESPYVEFPEPLTGYPG----- 350
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 303 PEAQ--RRVPKNPKYNTQRAERETQDSGSGNDGSGSEWEAQRDHLGPHRSTPESRAA 360
Qy 350 -EEL-----EEERGVKLGIDGFIYSVLVGAATAATGSDWNTTILACFVAILGLCL 401
Dy :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 361 VOELSGSILTSEDPEERGKVLGLGDFIFYSVLVGAATAATGSDWNTTILACFVAILGLCL 420
Qy 402 TLLILAIFKKALPALPISITFGLIFYSTDNLVRPFMDTLASHOLYI 448
Dy |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
Dy 421 TULLLAIFKKALPALPISITFGLVFFATDVLVQPFMDQLAFHQFYI 467

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RA ROYTA M., LILIUS L., EEROLA A., ST GEORGE-HYSLOP P.H., FREY H.,
RA LANNFELT L.;
RT "The Glu318Gly mutation of the presenilin-1 gene does not necessarily
RT cause Alzheimer's disease";
RL Ann. Neurol. 44:965-967(1998).
RN [15]
RP VARIANT GLY-318.
RX MEDLINE; 99066775.
RA ALDUDO J., BULLIDO M.J., FRANK A., VALDIVIESO F.;
RT "Missense mutation E318G of the presenilin-1 gene appears to be a
RT nonpathogenic polymorphism";
RL Ann. Neurol. 44:985-986(1998).
RN [16]
RP VARIANTS AD VAL-79; CYS-115; VAL-231, AND VARIANT GLY-318.
RX MEDLINE; 98046005.
RA CRUTS M., VAN DOIJN C.M., BROCKHOVEN H., VAN DEN BROECK M.,
RA WEHNET A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J.,
RA ST GEORGE-HYSLOP P.H., HOFMAN A., VAN BROECKHOVEN C.;
RT "Estimation of the genetic contribution of presenilin-1 and -2
RT mutations in a population-based study of presenile Alzheimer
RT disease";
RL Hum. Mol. Genet. 7:43-51(1998).
RN [17]
RP VARIANTS AD D-120; R-163; V-209; V-260; L-264; Y-410 AND P-426.
RX MEDLINE; 98180720.
RA POORAJ P., SHARMA V., ANDERSON L., NEMENS E., ALONSO M.E., ORR H.,
RA WHITE J., HESTON L., BIRD T.D., SCHELEBERG G.D.;
RT "Missense mutations in the chromosome 14 familial Alzheimer's disease
RT presenilin 1 gene";
RL Hum. Mutat. 11:216-221(1998).
RN [18]
RP VARIANT AD GLU-378.
RX MEDLINE; 99211215.
RA BESANCON R., LORENZI A., CRUTS M., RADAWIEC S., STURTZ F.,
RA BROUSSOLLE E., CHAZOT G., VAN BROECKHOVEN C., CHAMBA G.,
RA VANDENBERGHE A.;
RT "Missense mutation in exon 11 (codon 378) of the presenilin-1 gene in
RT a French family with early-onset Alzheimer's disease and transmission
RT study by mismatch enhanced allele specific amplification";
RL Hum. Mutat. 11:481-481(1998).
RN [19]
RP VARIANTS AD LEU-169 AND GLN-436.
RX MEDLINE; 99047368.
RA TADDEI K., KWOK J.B., KRIL J.J., HALLIDAY G.M., CREASEY H.,
RA HALLUPP M., FISHER C., BROOKS W.S., CHUNG C., ANDREWS C.,
RA MASTERS C.L., SCHOFIELD P.R., MARTINS R.N.;
RT "Two novel presenilin-1 mutations (Ser169Leu and Pro436Gln) associated
RT with very early onset Alzheimer's disease";
RL NeuroReport 9:3335-3339(1998).
RN [20]
RP VARIANT AD PRO-169.
RX MEDLINE; 99148656.
RA EZQUEPERRA M., CARNERO C., BLESER R., GELPI J.L., BALLESTA F., OLIVA R.;
RT "A presenilin 1 mutation (Ser169Pro) associated with early-onset AD
RT and myoclonic seizures";
RL Neurology 52:566-570(1999).

Query Match 62.8%; Score 1467; DB 1; Length 467;
Best Local Similarity 65.7%; Pred. No. 3.5e-94;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;

QY 24 ESTPTSCQGRGCPDGEANTAWRSQENDEEDGDDPRVCVSGVP-----GRPPG---- 75
DB 3 ELPAFLSYFQAOQMSDENHLSNTVRSQDNREOHNDNR-SLGHPEPLNGRPOGNSRQ 61
QY 75 -----LEELTLKYGAKVIMLFVPTLCMIVVATIKSVFYTEKNGQLIYPTPTD 127
DB 62 VFGQDEEDELTKYGAKVIMLFVPTLCMIVVATIKSVFYTEKNGQLIYPTPTD 121
QY 128 TPGVQRLNLSLNTLMISVIVVMTFLVLYKYRCYKFIHGWLIMSSLMLLFLFTYII 187
DB 122 TEIVGQRLNLSLNTLMISVIVVMTFLVLYKYRCYKFIHGWLIMSSLMLLFLFTYII 181

QY 188 LGEVLKTYNVAMDYPTLLLTWNFGAVGVMCIHWKGPLVLOQAYLIMISALMALVFIKYL 247
DB 182 LGEVFKTYNVADYITVALLIWNFGVGMISHWKGPLRLOQAYLIMISALMALVFIKYL 241
QY 248 PEWSAWVILGAIISVYDLVAVLCPKGPLRMLVETAOERNEPIFPALIISSAMVTVGMAKL 307
DB 242 PEWTAWLILAVISYDLVAVLCPKGPLRMLVETAOERNETLFPALIISSAMVTVGMVNAEG 301
QY 308 DPSSQGL--QLPYDPE-MEEDSDYDSFGE---PSYPEVFEPPLTGPQ----- 350
DB 302 DPEAQREVSKNSKYNASTERSQDTVAENDGGFSEWEAQRDSHLGPHRSTPESRAAV 361
QY 350 EEL-----EEEEERGKVLGLGDFIYFVSLVGRKAAATGSGDWNNTTACFAIILIGLCT 402
DB 362 QELSSSILAGEDPEERGKVLGLGDFIYFVSLVGRASATASGDWNNTTACFAIILIGLCT 421
QY 403 LLLAVFKKALPALPISITITGLIFSTDLNLRPFMDTLASHQLYI 448
DB 422 LLLAIFKKALPALPISITITGLIFSTDLNLRPFMDTLASHQLYI 467

RESULT 8
PSNL_RAT STANDARD; PRT; 468 AA.
AC P97887: P97529;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DE PRESENILIN 1 (PS-1) (S182 PROTEIN).
GN PSEN1 OR PSNL1.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-WISTAR: TISSUE-BRAIN;
RX MEDLINE; 97199371.
RA TANIGUCHI T., HASHIMOTO T., TANIGUCHI R., SHIMADA K., KAWAMATA T.,
RA YASUDA M., NAKAI M., TERASHIMA A., KOIZUMI T., MAEDA K., TANAKA C.;
RT "Cloning of the cDNA encoding rat presenilin-1";
RL Gene 186:73-75(1997).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-WISTAR: TISSUE-BRAIN;
RX MEDLINE; 9625262.
RA TAKAHASHI H., MURAYAMA M., TAKASHIMA A., MERCKEN M., NAKAZATO Y.,
RA NOGUCHI K., IMAHORI K.;
RT "Molecular cloning and expression of the rat homologue of
RT presenilin-1";
RL Neurosci. Lett. 206:113-116(1996).
CC -!- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE
CC EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE.
CC MAY FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS.
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC -----
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CC -----
DR EMBL; D82578; BAA11575.1; -;
DR EMBL; D82363; BAA11564.1; -;
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Glycoprotein.
FT TRANSWMEM 83 103 POTENTIAL.
FT TRANSWMEM 133 153 POTENTIAL.
FT TRANSWMEM 161 181 POTENTIAL.
FT TRANSWMEM 195 215 POTENTIAL.
FT TRANSWMEM 221 241 POTENTIAL.

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PSN1_MICMU STANDARD; PRT; 467 AA.
ID PSN1_MICMU STANDARD; PRT; 467 AA.
AC P79802;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN 1 (PS-1).
GN PSN1 OR PSN1 OR PS1.
OS Microcebus murinus (Lesser mouse lemur).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Strepsirrhini; Cheirogaleidae; Microcebus.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-BRAIN;
RX MEDLINE: 97079199.
RA CALEDA A., MESTRE-FRANCES N., CZECH C., PRADIER L., BONS N.,
RA BELLIS M.;
RT "Molecular cloning, sequencing, and brain expression of the
RT presentin 1 gene in Microcebus murinus.";
RL Biochem. Biophys. Res. Commun. 228:430-439(1996).
CC -1- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE
CC EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE.
CC MAY FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -1- ALTERNATIVE PRODUCTS: TWO ISOFORMS: I-467 (SHOWN HERE) AND I-
CC 463; ARE PRODUCED BY ALTERNATIVE SPLICING.
CC -1- TISSUE SPECIFICITY: FOUND PREDOMINANTLY IN NEURONS OF THE
CC DIFFERENT CORTICAL LAYERS AND HIPPOCAMPUS BUT ALSO IN SUBCORTICAL
CC STRUCTURES.
CC -1- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
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CC -----
DR EMBL; Z71333; CA95930.1; -
DR PFAM; PF01080; Presentinlin; 1.
KW Transmembrane; Alternative splicing; Glycoprotein.
FT TRANSMEM 83 103 POTENTIAL.
FT TRANSMEM 133 153 POTENTIAL.
FT TRANSMEM 161 181 POTENTIAL.
FT TRANSMEM 191 211 POTENTIAL.
FT TRANSMEM 221 241 POTENTIAL.
FT TRANSMEM 244 264 POTENTIAL.
FT TRANSMEM 281 301 POTENTIAL.
FT TRANSMEM 408 428 POTENTIAL.
FT TRANSMEM 433 453 POTENTIAL.
FT CARBOHYD 279 279 POTENTIAL.
FT CARBOHYD 405 405 POTENTIAL.
FT VARSPLIC 26 29 MISSING (IN ISOFORM I-463).
SQ SEQUENCE 467 AA; 52384 MW; A841A0B7 CRC32;

Query Match 61.9%; Score 1447; DB 1; Length 467;
Best Local Similarity 64.4%; Pred. No. 8.4e-93;
Matches 300; Conservative 41; Mismatches 83; Indels 42; Gaps 7;

QY 24 ESPTPRSCGRQGPGEQNTAQRWSENEEDGEDDDPRYVCSGVP-----GRPPG----- 75
Db 3 ELPAFLSYFQAQMSDNLHLSNTVRQNDREQQDGHGDRRL-GNPEPLSGRQPGNSGP 61
QY 75 -----LEELTKYGAHVIMLFPVTCMLVIVVATIKSVRYTEKNGOLIYTPPTED 127
Db 62 VVERDEEDELTKYGAHVIMLFPVTCMLVIVVATIKSVRYTRKDGOLIYTPPTED 121
QY 128 TPSVQGLLSNLTLTMSIVVMTIFLVLYKYRCYKFKHGLWLSLMLFLFTYIY 187
Db 122 TETVQGRALHSVLNAATMSIVVMTILLVLYKYRCYKVIHAWLISSLLLLFFFSFIY 181
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```
QY 188 LGEVLKTYNVAMDYPTLLLTWNFGAVGMVCIHWKGPLVLQOAYLIMISALMALVFIKYL 247
Db 182 LGEVFKTYNVAVDYITVALLTWNFGVGMISIHKGPLRLQOAYLIMISALMALVFIKYL 241
QY 248 PWSAWILGAIISYDVLVAVLCPKGPLRMLVETAOERNEPIFALIYSAMVYTMGMAKL 307
Db 242 FEWTAWLILAVISYDVLVAVLCPKGPLRMLVETAOERNEPIFALIYSSTMYLVNMAEG 301
QY 308 DPSQOGAL--OLPYD-----PEMEEDSDVSFGEPSYPEVPEP-----PLTGYPG 349
Db 302 DPEAQRVSKNTKYNAOQTEREAQASVPENDDGGFSEWEAQDSQLGPHRSTSVSRAAV 361
QY 350 EEL-----EEERGVKLGDFIFYSVLVGRKAAATGSGDWNNTLACFVAILIGLCIT 402
Db 362 QEISSSPASEDPEERGKVLGDFVYFVSVLVGRKASATASGDWNNTIACFVAILIGLCIT 421
QY 403 LLLLAFFKALPALPISITIFGLIFSTFNDLVPRPMDTILASHQLYI 448
Db 422 LLLLAIFKALPALPISITIFGLVFFATDYLQVPMQDLAFHOFYI 467

RESULT 11
PSN1_DROME STANDARD; PRT; 541 AA.
ID PSN1_DROME STANDARD; PRT; 541 AA.
AC O02194; O02395; O76802;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN HOMOLOG (DPS) (DMPS).
GN PS.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RX SEQUENCE FROM N.A. (LONG ISOFORM).
RX MEDLINE: 97285868.
RA BOULIANNE G.L., LYNE-BAR I., HUMPHREYS J.M., LIANG Y., LIN C.,
RA ROGAEV E., ST GEORGE-HYSLOP P.;
RT "Cloning and characterization of the Drosophila presentin
RT homologue.";
RL NeuroReport 8:1025-1029(1997).
RN [2]
RP SEQUENCE FROM N.A. (SHORT ISOFORM).
RC STRAIN-CANTON-S;
RA HONG C.S., KOO E.H.;
RT "Isolation and characterization of Drosophila presentin homolog.";
RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A. (LONG AND SHORT ISOFORMS).
RA YE Y., FORTINI M.E.;
RT "Characterization of Drosophila presentin and its colocalization
RT with Notch during development.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC -----
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CC -----
DR EMBL; U77934; AAB61139.1; -
DR EMBL; U78084; AAB53369.1; -
DR EMBL; AF084184; AAC33129.1; -
DR EMBL; AF084184; AAC33128.1; -
DR FLYBASE; FBgn0019947; PS.
DR PFAM; PF01080; Presentinlin; 1.
KW Transmembrane; Glycoprotein; Alternative splicing.
FT TRANSMEM 107 127
```

QY	324	EEDSYDSF-----GE-----PSYPEVFEP--	-----LTGYPGEEL-----	354
Db	279	EPTSSDNTSTAFPGASCSSSTPKRPKVRIPQKVQIESNTTASTQTSGVVRERELAA	338	
QY	354	-----EEERGKVLGLGFIFYSVLVGRAATGSDWNTTACFAVAILIGLC	400	
Db	339	ERTVDQANFHREEREERGKVLGLGFIFYSVLVGRASS--YFDWNTTIACYVAAILIGLC	396	
QY	401	LTLALLAVFKKALPALPISITFLIFVFFSTDN	432	
Db	397	FTLVLAVFKRALPALQ-----PFSPDS	420	
 RESULT 14				
HOPI-CAEEL				
AD	ID	HOPI-CAEEL	STANDARD;	PRT; 358 AA.
IC	AO	O02100;		
DT	15-JUL-1999	(Rel. 38, Created)		
DT	15-JUL-1999	(Rel. 38, Last sequence update)		
DT	15-DEC-1999	(Rel. 39, Last annotation update)		
DE	INTEGRAL MEMBRANE PROTEIN HOP-1.			
GN	HOP-1 OR C18E3.8.			
OS	Caenorhabditis elegans.			
OC	Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditiia; Rhabditida;			
OC	Rhabditia; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.			
RN	[1]			
RC	SEQUENCE FROM N.A.			
RP	STRAIN=BRISTOL N2;			
RC	MEDLINE: 98004548;			
RA	LI X., GREENWALD I.;			
RT	"HOP-1, a Caenorhabditis elegans presenilin, appears to be			
RT	functionally redundant with SEL-12 presenilin and to facilitate LIN-1			
RL	and GLP-1 signaling.";			
RN	Proc. Natl. Acad. Sci. U.S.A. 94:12204-12209(1997).			
CC	[2]			
RC	SEQUENCE FROM N.A.			
RP	STRAIN=BRISTOL N2;			
RA	CONNELL M., MAGGI L.;			
RL	Submitted (JUN-1997) to the EMBL/GenBank/DDBJ databases.			
CC	-!- FUNCTION: MAY FACILITATE LIN-12 AND GLP-1 MEDIATED RECEPTION OF			
CC	INTERCELLULAR SIGNALS.			
CC	-!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).			
CC	-!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.			
CC	-----			
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CC	or send an email to licenses@isb-sib.ch).			

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CC or send an email to license@isb-sib.ch).
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CC
CC EMBL; AF021905; AAB84394.1; -
CC EMBL; AF000265; AAB52948.1; -
CC WORMPEP; C18E3.8; CE08317.
CC PFAM; PF01080; Presenilin; 1.
KW Transmembrane.
FT TRANSMEM 13 33 POTENTIAL.
FT TRANSMEM 58 78 POTENTIAL.
FT TRANSMEM 87 107 POTENTIAL.
FT TRANSMEM 116 136 POTENTIAL.
FT TRANSMEM 149 169 POTENTIAL.
FT TRANSMEM 171 191 POTENTIAL.
FT TRANSMEM 298 318 POTENTIAL.
FT TRANSMEM 322 342 POTENTIAL.
SQ SEQUENCE 358 AA; 39864 MW; FC6305ED CRC32;
Query Match 22.5%; Score 524.5; DB 1; Length 358;
Best Local Similarity 30.5%; Pred. No. 2.1e-29;
Matches 117; Conservative 81; Mismatches 136; Indels 49; Gaps 8;
QY 83 YGAKHVIMLEVPVTLNCMIVVVAIKSVRYFTEKNGQLIYTPF-----TEDTPSVGQRLINS 138

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GenCore version 4.5
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OM protein - protein search, using sw model
Run on: March 18, 2000, 22:05:40 ; Search time 49.26 seconds
(without alignments)
630.566 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFMADSEEEVCDERTSL.....STDNLVRPFMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 225878 seqs, 69334122 residues

Database : SPTREMBL_12.*

Word size : 0

Number of hits that pass the threshold : 225878

- 1: sp_archaea.*
- 2: sp_bacteria.*
- 3: sp_fungi.*
- 4: sp_human.*
- 5: sp_invertebrate.*
- 6: sp_mammal.*
- 7: sp_mhc.*
- 8: sp_organelle.*
- 9: sp_phage.*
- 10: sp_plant.*
- 11: sp_rodent.*
- 12: sp_virus.*
- 13: sp_vertebrate.*
- 14: sp_unclassified.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	2262.5	96.9	449	6 Q9XT96	Q9xt96 bos taurus
2	1438.5	61.6	478	6 Q9XT97	Q9xt97 bos taurus
3	1407.5	60.3	456	13 Q9W6T7	Q9w6t7 brachydanio
4	1355	58.0	384	13 Q73869	Q73869 cyprinus ca
5	576.5	24.7	272	5 Q96340	Q96340 drosophila
6	440	18.8	184	4 Q95465	Q95465 homo sapien
7	113.5	4.9	406	5 Q19737	Q19737 caenorhabdi
8	112.5	4.8	4578	13 Q42181	Q42181 fugu rubrip
9	110.5	4.7	320	8 Q34086	Q34086 coccyzus er
10	110.5	4.7	381	8 Q35425	Q35425 phascosolor
11	109	4.7	1840	11 Q70611	Q70611 rattus norv
12	108.5	4.6	380	8 Q32209	Q9zcc9 upupa epops
13	107.5	4.6	748	2 Q32577	Q9z577 streptomyce
14	107	4.6	381	8 Q33723	Q33723 antechinus
15	106.5	4.6	382	8 Q34340	Q34340 didelphis m
16	106.5	4.6	318	11 Q35294	Q35294 rattus norv
17	106.5	4.6	303	11 P97829	P97829 rattus norv
18	106	4.5	652	5 Q93346	Q93346 caenorhabdi
19	106	4.5	382	8 Q34279	Q34279 didelphis a
20	106	4.5	382	8 Q34677	Q34677 glirolia ve
21	106	4.5	1717	13 Q90519	Q90519 fugu rubrip
22	105	4.5	444	2 Q9X2N3	Q9x2n3 arthrobacte
23	105	4.5	379	8 Q34428	Q34428 echimys did
24	104	4.5	382	8 Q35561	Q35561 philander o
25	104	4.5	379	8 Q36096	Q36096 trinomys pa

ALIGNMENTS

```
RESULT 1
Q9XT96
ID Q9XT96 PRELIMINARY; PRT; 449 AA.
AC Q9XT96;
DT 01-NOV-1999 (Tremblrel. 12, Created)
DT 01-NOV-1999 (Tremblrel. 12, Last sequence update)
DT 01-NOV-1999 (Tremblrel. 12, Last annotation update)
DE PRESENILIN 2.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovinae; Bos.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=BRAIN;
RA SAHARA N., SHIRASAWA T., MORI H.;
RT "Molecular cloning of bovine presenilin 2 gene.";
RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF038937; AAD39024.1; -
SQ SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;
```

Query Match	96.9%	Score 2262.5;	DB 6;	Length 449;
Best Local Similarity	97.1%	Pred. NO. 4.7e-162;		
Matches 436;	Conservative 5;	Mismatches 7;	Indels 1;	Gaps 1;
QY 1	MLTFMADSEEEVCDERTSLMSAESPTPRSCQGRGPDGENTAWRSQENEED-GEED	59		
Db 1	MLTFMADSEEEVCDERTSLMSAESPTPRSCQGRGLEDGESAAQWRSQESDEHDEED	60		
QY 60	PRYVCVGVPGRPPGLEELTLKYAKHVMFLFVPTVLCMVVVVATIKSVRFYTEKNGQL	119		
Db 61	PRYVCVGVPGRPPGLEELTLKYAKHVMFLFVPTVLCMVVVVATIKSVRFYTEKNGQL	120		
QY 120	ITYPTFEDPSPVGORLLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGLIMSSLM	179		
Db 121	ITYPTFEDPSPVGORLLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGLIMSSLM	180		
QY 180	LELFTYIYGEVLKYNVAMDYPTLLTYVNFAGVMCIHWKGPLVLOQAYLIMISALM	239		
Db 181	LELFTYIYGEVLKYNVAMDYPTLLTYVNFAGVMCIHWKGPLVLOQAYLIMISALM	240		
QY 240	ALVFIKYLPEWSAWTILGAISVYDLVAVLCPKGPLRMLVETAQERNEPIFALIYSSAMV	299		
Db 241	ALVFIKYLPEWSAWTILGAISVYDLVAVLCPKGPLRMLVETAQERNEPIFALIYSSAMV	300		
QY 300	WTVGMAKLPDPSQALQLPYDPMEEDSDSDSFCGEPSPYFVFEPPLTGTGPGEELEBERG	359		

FN SEQUENCE FROM N.A.
RA ARCHER S., HIRANO J., DISS J.K., FRASER S.P., DJAMGOZ M.B.A.;
RL NeuroReport 0:0-0(0).
DR EMBL; Y17128; CAA76641.1; -.
RR PFAM; YF01080; Presentinlin; 1.
FT NON_TER 1
SQ SEQUENCE 384 AA; 43276 MW; 21A78D17 CRC32;

Query Match 58.0%; Score 1355; DB 13; Length 384;
Best Local Similarity 69.9%; Pred. No. 3.9e-94;
Matches 270; Conservative 33; Mismatches 55; Indels 28; Gaps 5;

Qy 89 IMLFVPVTLGVVATKSVRFYTEKNG-QLIYTPFTEDTSPVQRLNLSVNTLMIS 147
Db 1 IMLFIPVTLGVVATKSVFYTKDQGLIYTPFREDTETVQRLNLSMLNAIMIS 60
Qy 148 VIVVMTIFLVLYKYRCYKEIHGWLIMSLMLLFTYIYLGEVLKTYNVANDYPTLLT 207
Db 61 VIVVMTIFLVLYKYRCYKVIQGWLFNLLFFSFIYLGVEFKTYNVANDYPTLAVI 120
Qy 208 VNFAGVGVCIHMKGPLVLAQAYLIMISALMALVFIKYLPEWSAWVILGALSVDLVAV 267
Db 121 IWNFGVGVCIHMKGPLRLQAYLIMISALMALVFIKYLPEWTAWLILAAISVYDLLAV 180
Qy 268 LCPKGPLRLVETAEQNEPIFPALIIYSAMVTVGMA-KLDPSSQGLALQLP----- 319
Db 181 LCPKGPLRLVETAEQNEPIFPALIIYSAMVTVGMA-KLDPSSQGLALQLP----- 319
Qy 319 -----YDPEMEEDSYDSFG-----EPSYVEPEPLTGYTGELEEEERGVL 362
Db 241 APTAQPEDGGFTFANVQQHQGLQPMOSTEDSRREIOLPSARPP--PVEDDEERGVL 298
Qy 363 GLGDFIFYSVLVGRKAATGSDGNTTTLACFVAILIGLCTLLLLAVFKKALPALPISITF 422
Db 299 GLGDFIFYSVLVGRKASATGSDGNTTTLACFVAILIGLCTLLLLAVFKKALPALPISITF 358
Qy 423 GLIFVFSTDLVRRPMDPLASHQLYI 448
Db 359 GLVFYFATDLVRRPMDQLAVHQFYI 384

RESULT 5
ID O96340 PRELIMINARY; PRT; 272 AA.
AC O96340;
DT 01-MAY-1999 (TremBLrel. 10, Created)
DT 01-MAY-1999 (TremBLrel. 10, Last sequence update)
DT 01-MAY-1999 (TremBLrel. 10, Last annotation update)
DE PRESENILIN (FRAGMENT).
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CANTON S;
RX MEDLINE; 98331525.
RA MAREFANY G., DEL-FAVERO J., VALERO R., DE JONGHE C., WOODROW S.,
RA HENDRIKS L., VAN BROECKHOVEN C., GONZALEZ-DUARTE R.;
RT "Identification of a Drosophila presentinlin homologue: evidence of
RT alternatively spliced forms";
RL J. Neurogenet. 12:41-54(1998).
DR EMBL; AF017025; AAD01611.1; -.
FT NON_TER 1
SQ SEQUENCE 272 AA; 29456 MW; 606B9A5C CRC32;

Query Match 24.7%; Score 576.5; DB 5; Length 272;
Best Local Similarity 47.8%; Pred. No. 5.7e-36;
Matches 133; Conservative 29; Mismatches 41; Indels 75; Gaps 8;

Qy 240 ALVFIKYLPEWSAWVILGALSVDLVAVLCPKGPLRLVETAEQNEPIFPALIIYSAMV 299
Db 1 ALVFIKYLPEWTANAVLAASISWDLAVLSRPGPLRLVETAEQNEPIFPALIIYSTVV 60
Qy 300 WTV-----GMAKLDPPS-----QGALQLPYDPEMEEDSYDSFGPEPSYE- 339
Db 61 YALVNTVTPQQSOATASSPSSSNSTTTTTRATQNSLA---SPEAAAAAGORTGN-SHPRQ 116
Qy 339 -----VFEPPLTGYGCE---ELEE----- 355
Db 117 NORDDCSVLATEAEAAAGFTQEWNSANLSERVARRQIEVQSTQSGNAQORSNEYRTVTPADQN 176
Qy 355 -----EEERGKVLGDFIYFSLVGRKAATGSDGNTTTLACFVAILIGLCTLLLLAVFK 410
Db 177 HPDQGEERIKLGLGDFIYFSLVGRKASS--YGDWTTTACFVAILIGLCTLLLLAIWR 234
Qy 411 KALPALPISITITGLIFGFYFSTDMNLVRRPMDPLASHQLYI 448
Db 235 KALPALPISITITGLIFCFATSAVVKPFMEDLSAKQVFI 272

RESULT 6
ID O95465 PRELIMINARY; PRT; 184 AA.
AC O95465;
DT 01-MAY-1999 (TremBLrel. 10, Created)
DT 01-MAY-1999 (TremBLrel. 10, Last sequence update)
DT 01-MAY-1999 (TremBLrel. 10, Last annotation update)
DE MINILIN.
GN PSNI.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RA POWELL C.S., GEGG M.E., PALMER M.S.;
RT "Human presentinlin 1 gene encodes an alternative protein-minilin.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ008005; CAA07825.1; -.
SQ SEQUENCE 184 AA; 21073 MW; 5C6FBAEE CRC32;

Query Match 18.8%; Score 440; DB 4; Length 184;
Best Local Similarity 56.0%; Pred. No. 6.2e-26;
Matches 102; Conservative 15; Mismatches 43; Indels 22; Gaps 4;

Qy 24 ESPTPRSCOEGRQGEDGENTAQWRSQENEDEGEDDPDRYVCSGVP-----GRPPG--- 75
Db 3 ELPAPLSYFQNAQMSDNLSTNTRVSDNDRERQENDRR-SLGHPEPLSNGRPQNSRQ 61
Qy 75 -----LEEELTKYGAHVIMLFVPTLCMIVVATIKSVFYTEKNGQLIYTPFTED 127
Db 62 VVEQDEEEDDEULTKYGAHVIMLFVPTLCMIVVATIKSVFYTRKDGQLIYTPFTED 121
Qy 128 TFSVQRLNLSVNTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSMLLFLFTYIY 187
Db 122 TETVQRLHSLNLAIMISVIVMTIFLVLYKYRCYK-----VSMRHSLLSTLFFLW 176
Qy 188 LG 189
Db 177 LG 178

RESULT 7
ID Q19737 PRELIMINARY; PRT; 406 AA.
AC Q19737; Q22692;
DT 01-NOV-1996 (TremBLrel. 01, Created)
DT 01-MAY-1999 (TremBLrel. 10, Last sequence update)
DT 01-NOV-1999 (TremBLrel. 12, Last annotation update)
DE F22E10.5 PROTEIN.
GN F22E10.5.
OS Caenorhabditis elegans.

Qy	211	FGAVGVCVHWKGPLVLQQAAYLIMISALMALVP	IKYLP	PEWSAWVILGALSV	VDLVAVLCP	270	
Db	115	IG-----	VILLAVATAFGVGLPWQ	SEFWGATVITNL	LSAIPY	155	
Qy	271	KGPLRLVE-----	TAQERNEP	IPFALLIYSSAMVY	VTGMAXL	312	
Db	156	IGP-"ILA	EWVWG	GYADV	KATLRFPAFH	FLPFIVTALAIVHLLF	113
Qy	313	GALQLYDPD-----	MEESYDS	FGSEPS	YEPVEP	352	
Db	214	NADKIPFHYPTIKDALG	FMLLSVLLLT	LFSPD	SLGD---	PDNFS	270
Qy	353	EEEEERGVLGLG	DFIF-YSVLV	GKAAATGSGD	WNMTLAC	FVAILIGLIGLTL	411
Db	271	E-----	WYFIF	FAVAILRS	INKLGG----	VLLASILILLIPL	313
Qy	412	ALPALPISITFGLIF	YFST	DNLV	434		
Db	314	SMF	FRISOT---	LFWILT	ANLI	333	

RESULT 11	
ID 070611	
ID 070611	PRELIMINARY; PRT: 1840 AA.
AC 070611;	
DT 01-AUG-1998	(TREMBLrel. 07, Created)
DT 01-AUG-1998	(TREMBLrel. 07, Last sequence update)
DT 01-NOV-1998	(TREMBLrel. 07, Last annotation update)
DT 01-NOV-1998	(TREMBLrel. 12, Last annotation update)
DE	RAT SKELETAL MUSCLE TYPE 1 VOLTAGE-GATED SODIUM CHANNEL
DE	(R8KM1) VARIANT (R8KM1).

DE (RSKML) VARIANI (RSKML).
GN SCN4A.
OC Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]
RN SEQUENCE FROM N.A.
RP STRAIN=COPENHAGEN; TISSUE=PROSTATE;
RC DISS J.K., STEWART D., BLACK J.A., FRASER S.P., DIBB-HAJJ S.,
RA WAXMAN S.G., ARCHER S., DJANGOZ M.B.A.;
RL FEBS Lett. 0:0-0(0).
DR EMBL; Y17153; CAA76659.1; -.
DR PFAM; PF00520; ion.trans. 4.
DR PFAM; PF00612; IQ; 1.
DR PRINTS; PR00170; NACHANNEL.
DR Ionic channel.
SQ SEQUENCE 1840 AA; 208823 MW; 1948B0C2 CRC32;

Query Match 4.7%; Score 109; DB 11; Length 1840;
Best Local Similarity 20.3%; Pred. No. 5.4;
Matches 98; Conservative 63; Mismatches 130; Indels 192; Gaps

```

QY      5 MASDSEEEVDERTSLM-----SASPTPRSCOEGRQGP-E-DGENTAQRSENEE---55
       :|::||::||::||::||::||::||::||::||::||::||::||::||::||:
Db     452 LAEDQSKE---EEFQOMLEKKYKHELEKAKAAQALESGBEADGDT-----HNKDNG 503
       :|::||::||::||::||::||::||::||::||::||::||::||::||:
QY     55 -----DGEEDPDRVCSGVGPGRPLGEE-----ELTIKYGA-----KH 87
       :|::||::||::||::||::||::||::||::||::||::||::||::||:
Db    504 SLDSASEKGPPRSCSAADS AISDA MELEFAHOKCPWPKCAHKVIWNCAAPWKFKH 563

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QY 88 VMLFV-----PVTLCMLVWVATIKSVRFYTEKNGQLIYTPTE-----DTPSVGQRLN 137
DB 564 IYILVMDPFVDLGITIC-IVNLTFMAMEHY-----PMTHEFDNVLSVGNLVFT 612

[illegible]

DT 01-MAY-1999 (TREMBLrel. 10, Last annotation update)
DE CYTOCHROME B.
OS Antechinus naso (Antechinus habbema).
OG Mitochondrion.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Metatheria; Dasyuromorphia; Dasyuridae; Antechinus.
RN [1]
RP SEQUENCE FROM N.A.
RA KRAJEWSKI C., PAINTER J., DRISKEILL A.C., BUCKLEY L., WESTERMAN M.;
RA Sci. New Guinea 19:157-166(1993).
RN [2]
RP SEQUENCE FROM N.A.
RA KRAJEWSKI C., PAINTER J., BUCKLEY L., WESTERMAN M.;
RA J. Mammal. Evol. 2:25-35(1994).
RN [3]
RP SEQUENCE OF 1-287 FROM N.A.
RA KRAJEWSKI C., BUCKLEY L., WOOLLEY P.A., WESTERMAN M.;
RA J. Mammal. Evol. 3:81-91(1996).
RN [4]
RP SEQUENCE FROM N.A.
RA ARMSTRONG L.A., KRAJEWSKI C., WESTERMAN M.;
RA J. Mammal. 0:0-0(1998).
RN [5]
RP SEQUENCE FROM N.A.
RA KRAJEWSKI C.;
RA Submitted (MAR-1994) to the EMBL/GenBank/DBJ databases.
RN [6]
RP SEQUENCE FROM N.A.
RA KRAJEWSKI C.;
RA Submitted (FEB-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; U07576; AAC03634.1; -;
DR PFAM; PF00032; cytochrome_b_C; 1.
DR PFAM; PF00033; cytochrome_b_N; 1.
KW Mitochondrion.
SQ SEQUENCE 381 AA; 42738 MW; 9C6998A0 CRC32;

Query Match 4.6%; Score 107; DB 8; Length 381;
Best Local Similarity 21.3%; Pred. No. 1.3;
Matches 76; Conservative 53; Mismatches 111; Indels 116; Gaps 17;

QY 138 SVLNTLMISVIVMTIFLVLY-----KYRCYKFTGWLIM-----SSLMLL 180
DB 138 SVLNTLMISVIVMTIFLVLY-----KYRCYKFTGWLIM-----SSLMLL 180
QY 181 FLFTYIYLGEVLTNYVNDYPTLLTYNFGAVGVCIHWRKGPLVLOQAYLIMISALMA 240
DB 181 FLFTYIYLGEVLTNYVNDYPTLLTYNFGAVGVCIHWRKGPLVLOQAYLIMISALMA 240
QY 241 LVFTKYLPEWSANVILGSAISYVDLVAVLCPKGPLRLMVE-----TAQERNEP 287
DB 241 LVFTKYLPEWSANVILGSAISYVDLVAVLCPKGPLRLMVE-----TAQERNEP 287
QY 288 IFPALIYSSAMVWTVGM-----AKLDPSSOGALQLPYDPE----- 323
DB 288 IFPALIYSSAMVWTVGM-----AKLDPSSOGALQLPYDPE----- 323
QY 323 -NEEDSDSFGEPSEYVEFEP--PLTGYPGEELEEEERGVKGLGDFIF--YSLVGVKAA 378
DB 323 -NEEDSDSFGEPSEYVEFEP--PLTGYPGEELEEEERGVKGLGDFIF--YSLVGVKAA 378
QY 379 ATGSGDNWTTLACFVAILIGLCLTLLAVFKKALPALPISITFGLIFFSTDNLV 434
DB 379 ATGSGDNWTTLACFVAILIGLCLTLLAVFKKALPALPISITFGLIFFSTDNLV 434
QY 409 FKKALPALPISITFGLIFFSTDNLV 434
DB 409 FKKALPALPISITFGLIFFSTDNLV 434

RESULT 15

Q34340 ID Q34340 PRELIMINARY; PRT; 382 AA.
AC Q34340;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-MAY-1999 (TREMBLrel. 10, Last annotation update)

DE CYTOCHROME B LIGHT STRAND.
OS Didelphis marsupialis.
OG Mitochondrion.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Metatheria; Didelphimorphia; Didelphidae; Didelphis.
RN [1]
RP SEQUENCE FROM N.A.
RA PATON J.L., REIS MARIA S.F., DA SILVA N.F.;
RA J. Mammal. Evol. 3:3-29(1996).
RN [2]
RP CATALYTIC ACTIVITY: OH(2) + 2 FERRICYTOCHROME C = Q + 2
CC FERROCYTOCHROME C.
CC -!- COFACTOR: TWO HEME GROUPS (B562 AND B566) WHICH ARE NOT COVALENTLY
CC BOUND TO THE PROTEIN (BY SIMILARITY).
DR EMBL; U34655; AAA99746.1; -;
DR PFAM; PF00032; cytochrome_b_C; 1.
DR PFAM; PF00033; cytochrome_b_N; 1.
KW Mitochondrion; Electron transport; Respiratory chain; Transmembrane;
Heme.
SQ SEQUENCE 382 AA; 43139 MW; 907FBCA3 CRC32;

Query Match 4.6%; Score 106.5; DB 8; Length 382;
Best Local Similarity 20.5%; Pred. No. 1.4;
Matches 79; Conservative 53; Mismatches 115; Indels 139; Gaps 18;

QY 97 LCMIVVATIKSVRYTEKNGQLIYPTFTEDTPSGQRLNLSVNLIMLSIVVMTIFL 156
DB 97 LCMIVVATIKSVRYTEKNGQLIYPTFTEDTPSGQRLNLSVNLIMLSIVVMTIFL 156
QY 157 VVLYKYRCYKFTGWLIM-----SSLMLEFLFTYIYLGEVLTNYVNDYPTLLTVMN 210
DB 157 VVLYKYRCYKFTGWLIM-----SSLMLEFLFTYIYLGEVLTNYVNDYPTLLTVMN 210
QY 65 SV--AHICRDVNYGLIRNIHANGASMFCLFLHVGRIYGY-----LYKETWN 114
DB 65 SV--AHICRDVNYGLIRNIHANGASMFCLFLHVGRIYGY-----LYKETWN 114
QY 211 FGAVGVNCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSANVILGSAISYVDLVAVLCP 270
DB 211 FGAVGVNCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSANVILGSAISYVDLVAVLCP 270
QY 115 IG-----VILLTVMATAFVGVLPGQMSFWGATVITNLSAIPY 155
DB 115 IG-----VILLTVMATAFVGVLPGQMSFWGATVITNLSAIPY 155
QY 271 KGPLRLMVE-----TAQERNPIPPALYSSAMVWTVGMK-----LDP 309
DB 271 KGPLRLMVE-----TAQERNPIPPALYSSAMVWTVGMK-----LDP 309
QY 156 IG--NTLVEWINGGFSVDKATLTTRFAFHFIPLFIILAMVWVHLLFTHETGNNPTGLDP 213
DB 156 IG--NTLVEWINGGFSVDKATLTTRFAFHFIPLFIILAMVWVHLLFTHETGNNPTGLDP 213
QY 310 SSOGALQLPYDPE-----EMEEDSYDSFGEPSEYVEFEP--PLTGYPG 349
DB 310 SSOGALQLPYDPE-----EMEEDSYDSFGEPSEYVEFEP--PLTGYPG 349
QY 214 NSD---KIPFHPYTYIKOILGLFLMTIILLSLAMFSPOLLGD---PDNFTPANPLNTPPH 267
DB 214 NSD---KIPFHPYTYIKOILGLFLMTIILLSLAMFSPOLLGD---PDNFTPANPLNTPPH 267
QY 350 ELEEEERGVKGLGDFIF--YSLVGVKAAATGSGDNWTTLACFVAILIGLCLTLLAV 408
DB 350 ELEEEERGVKGLGDFIF--YSLVGVKAAATGSGDNWTTLACFVAILIGLCLTLLAV 408
QY 268 IKPE-----WYFLFAYAILRSIPNKLGG-----VLALLASILILIMPLLTST 311
DB 268 IKPE-----WYFLFAYAILRSIPNKLGG-----VLALLASILILIMPLLTST 311
QY 409 FKKALPALPISITFGLIFFSTDNLV 434
DB 409 FKKALPALPISITFGLIFFSTDNLV 434

Search completed: March 18, 2000, 22:07:43
Job time: 123 sec

=> e

E6 1 PRESENILIN-2 PS2S (MOUSE PS-2SHORT ISOFORM)/CN
E7 1 PRESENILIN-ASSOCIATED PROTEIN 1 (HPAP-1) (HUMAN INCYTE
CLONE 1353337)/CN
E8 1 PRESENILINASE/CN
E9 1 PRESEP-AGRI/CN
E10 1 PRESER ACE/CN
E11 1 PRESERCAR/CN
E12 1 PRESERIN M 72/CN
E13 1 PRESERIN T 72/CN
E14 1 PRESERT/CN
E15 1 PRESERVAC WETPROOF/CN
E16 1 PRESERVAL/CN
E17 1 PRESERVAL B/CN

=> s e4-6

L1 1 "PRESENILIN-2 (HUMAN ISOFORM)"/CN
1 "PRESENILIN-2 PS2CCAS (MOUSE)"/CN
1 "PRESENILIN-2 PS2S (MOUSE PS-2SHORT ISOFORM)"/CN
3 ("PRESENILIN-2 (HUMAN ISOFORM)"/CN OR "PRESENILIN-2 PS2CCAS
(MOUSE)"/CN OR "PRESENILIN-2 PS2S (MOUSE PS-2SHORT
ISOFORM)"/CN)

=> d 1-3 ide can

L1 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2000 ACS
RN 251358-30-2 REGISTRY
CN **Presenilin-2 (human isoform) (9CI)** (CA INDEX NAME)
OTHER NAMES:
CN 5: PN: WO9960122 SEQID: 5 claimed protein
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
*** USE 'SQD' OR 'SQIDE' FORMATS TO DISPLAY SEQUENCE ***
1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 132:11416

L1 ANSWER 2 OF 3 REGISTRY COPYRIGHT 2000 ACS
RN 200445-64-3 REGISTRY
CN 330-448-presenilin-2 (Mus musculus isoform PS2Ccas) (9CI) (CA INDEX
NAME)
OTHER NAMES:
CN **Presenilin-2 PS2Ccas (mouse)**
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***